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<p>(21) International Application Number: PCT/GB90/00753</p> <p>(22) International Filing Date: 16 May 1990 (16.05.90)</p> <p>(30) Priority data:</p> <table> <tr><td>8911295.7</td><td>17 May 1989 (17.05.89)</td><td>GB</td></tr> <tr><td>8916725.8</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> <tr><td>8916726</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> <tr><td>8916727.4</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> <tr><td>8916728.2</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> <tr><td>8916729.0</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> <tr><td>8916730.8</td><td>21 July 1989 (21.07.89)</td><td>GB</td></tr> </table> <p>(71) Applicant (for all designated States except US): IMPERIAL CHEMICAL INDUSTRIES PLC [GB/GB]; Imperial Chemical House, Millbank, London SW1P 3JF (GB).</p> <p>(72) Inventors; and</p> <p>(75) Inventors/Applicants (for US only) : BRIGHT, Simon, William, Jonathan [GB/GB]; 24 Pound Lane, Marlow, Bucks SL7 2AY (GB). CHANG, Ming, Tang [CN/US]; 1419 Illinois Avenue, Ames, IA 50010 (US). EVANS, Ian, Jeffrey [GB/GB]; 38 Wilmington Close, Woodley, Reading, Berkshire RG5 4LR (GB). MACDONALD, Mairi, Jean [GB/GB]; 35 Flamingo Close, Wokingham, Berkshire RG11 2SJ (GB).</p>		8911295.7	17 May 1989 (17.05.89)	GB	8916725.8	21 July 1989 (21.07.89)	GB	8916726	21 July 1989 (21.07.89)	GB	8916727.4	21 July 1989 (21.07.89)	GB	8916728.2	21 July 1989 (21.07.89)	GB	8916729.0	21 July 1989 (21.07.89)	GB	8916730.8	21 July 1989 (21.07.89)	GB	<p>(74) Agent: HUSKISSON, Frank, Mackie; Imperial Chemical Industries plc, Legal Dept, Patents, PO Box No. 6, Bessemer Road, Welwyn Garden City, Herts AL7 1HD (GB).</p> <p>(81) Designated States: AT (European patent), AU, BB, BE (European patent), BF (OAPI patent), BG, BJ (OAPI patent), BR, CA, CF (OAPI patent), CG (OAPI patent), CH (European patent), CM (OAPI patent), DE (European patent)°, DK (European patent), ES (European patent), FI, FR (European patent), GA (OAPI patent), GB (European patent), HU, IT (European patent), JP, KP, KR, LK, LU (European patent), MC, MG, ML (OAPI patent), MR (OAPI patent), MW, NL (European patent), NO, RO, SD, SE (European patent), SN (OAPI patent), SU, TD (OAPI patent), TG (OAPI patent), US.</p> <p>Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>	
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<p>(54) Title: HERBICIDE RESISTANT MAIZE</p> <p>(57) Abstract</p> <p>Herbicide resistant maize plants of inbred line UE95 are obtained by mutagenesis of pollen followed by selection by germination of the resulting seed in the presence of the herbicide. The plants are resistant to normally lethal concentrations of the herbicide imazethapyr (PURSUIT) (American Cyanamid) and have resistance to other herbicides of the imidazolinone family and to sulphonylurea herbicides such as chlorsulfuron.</p>																								

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HERBICIDE RESISTANT MAIZE

This invention relates to herbicide resistant maize plants and the seed and progeny thereof.

The purpose in providing crop plants which resist the action of a herbicide is to facilitate the destruction of weeds growing between the plants by the overall application of a herbicidally effective concentration of a herbicide which would destroy the crop plant in its normal, that is herbicide sensitive, state. Such resistant plants are also useful for use in a locus of any short term carry-over of herbicide from a previous crop.

Methods are known by which populations of plants may be obtained which contain a great number of random mutations. Such methods include tissue culture techniques where spontaneous somaclonal variation occurs in the presence or absence of a mutagen. By applying to the cultures some form of selection pressure it is possible to recover cells which resist that pressure. Depending on the plant species it is sometimes possible to regenerate whole plants from the resistant cells. Such tissue culture selection methods have been used in the past to select for resistance to herbicides.

Neuffer and Coe [Maydica XXIII (1978) 21-28; page 21] have described a procedure for corn (maize) pollen mutagenesis using mutagens suspended in light paraffin oil, followed by pollination of a recipient plant with the mutagenized pollen. As reported, there is no indication that any attempt was ever made to locate and isolate commercially important mutants and, although it is said that the resulting plants

were examined for mutants, no details are given as to the properties of any mutants which may have been found, probably by visual inspection.

In Plant Breeding Reviews 5, pages 39 to 180, 5 Bird and Neuffer review the uses of mutagenic processes to produce variation in maize. Pollen mutagenesis is discussed at pages 150 and 151. Pollen mutagenesis gives a relatively high frequency of variation in the M_1 generation compared with other 10 available procedures. However, no suggestion is made in respect of the use of pollen mutagenesis for generation of mutants which display resistance to herbicide action. Section III (page 149) describes some of the difficulties of the use of mutagenesis as 15 a source of genetic variation. One important aspect is the method by which the population of putative mutants is screened for useful phenotypes.

An object of the present invention is to provide herbicide resistant maize.

20 According to the present invention there is provided a method of producing mutant Zea mays possessing resistance to a normally lethal concentration of a chosen herbicide, comprising subjecting isolated pollen of Zea mays to the action 25 of a chemical mutagen, recovering mutagenised pollen, pollinating a female parent therewith, harvesting seed from the mature plant, growing the seeds under selection pressure of the chosen herbicide or a second herbicide having the same mode of action as 30 the chosen herbicide, and recovering progeny resistant to the applied selection pressure.

Preferably the chemical mutagen is ethyl methanesulphonate.

Preferably the herbicide utilised for applying

selection pressure is an inhibitor of acetolactate synthase and more preferably is imazethapyr.

Preferably also the selection pressure is applied by application of the herbicide to the seeds 5 pre-emergence.

It is preferred that the concentration of herbicide used for selection pressure is insufficient to inhibit germination but sufficient to inhibit growth of sensitive plants.

10 Further according to the present invention we provide herbicide resistant maize plants, seeds of which have been deposited at the National Collection of Industrial & Marine Bacteria, Aberdeen, United Kingdom, official details of the deposits being as in 15 Table 1 below.

TABLE 1

Mutant Number	Applicant Code	Accession Number	Date of Deposit
20	1 ICI/88/01 63-3D	NCIMB 40137	8 May 1989
	2 ICI/88/03 56-3G	NCIMB 40136	8 May 1989
	3 ICI/88/04 82-5H	NCIMB 40138	8 May 1989
	4 see footnote		~
25	5 ICI/88/07 97-4L	NCIMB 40167	18 July 1989
	6 ICI/88/07 76-6H	NCIMB 40168	18 July 1989
	7 ICI/88/07 58-5D	NCIMB 40169	18 July 1989
	8 ICI/88/09 77-7N	NCIMB 40170	18 July 1989
	9 ICI/88/10 1-4C	NCIMB 40171	18 July 1989
	10 ICI/88/10 72-4N	NCIMB 40172	18 July 1989

30 * Mutant Number 4 was withdrawn in the light of

further testing which showed that it possessed no useful resistance. The number has been retained to retain the sequence on numbers in the description which follows.

5 Each of these deposits is a genetic mixture, segregating mutant and non-mutant seeds. The mutants are heterozygous for the gene conferring herbicide resistance. Mutant plants may be derived from these seeds by growing under the conditions described below
10 under the heading "Screening" in the presence of the herbicide imazethapyr.

The invention also provides seeds of the said plant and progeny of the said plant which have been produced by crossing of the plants of this invention
15 with other maize plant lines.

The plants of this invention are known to be
resistant to certain members of the imidazolinone
family of herbicides, for example, imazethapyr [5-
20 ethyl-2-(5-iso-propyl-5-methyl-4-oxo-2-imidazolin-
2-yl) nicotinic acid: (Trade Mark PURSUIT, American
Cyanamid)]. Cross resistance to herbicides of the
sulphonylurea family, chlorsulfuron, for example, has
also been found, as has resistance to the phenoxy-
25 pyrimidines and the triazolopyrimidines. Resistance
to other, as yet untested, herbicide families may also
be present.

The herbicides against which the plants of the
invention have so far been tested are those whose mode
of action is inhibition of the enzyme acetolactate
30 synthase (ALS), also known as acetohydroxyacid
synthase (AHAS). It is convenient to refer to the
herbicides by the Trade Marks by which they are sold
commercially. Figure 2 of the drawings gives the
chemical structures of the active ingredients.

A detailed exposition of the molecular basis of the resistance to the sulphonylurea herbicides is given in European Patent Application 257,993 (E.I. Du Pont de Nemours and Company). Isolation of 5 imidazolinone-resistant maize from tissue culture is reported in United States Patent Number 4,761,373 (Molecular Genetics Inc.). Various plant species and several herbicides are reported in the literature as having been used in tissue culture processes to 10 isolate resistant mutants.

In comparison, the present invention does not utilise tissue culture. The resistance is introduced by mutagenesis of pollen by a chemical mutagen. Selection is effected directly on the seeds formed 15 after fertilisation with the mutagenised pollen either by treatment of the seed pre-emergence or at the seedling stage, or both.

Certain advantages accrue from the use of pollen mutagenesis as a method of creating mutants rather 20 than the more usual method of relying on somaclonal variation to produce the variation. In order that somaclonal variation may occur it is required that a culture of plant tissue be established. This requirement restricts the choice of genotype which may 25 be used as it is not always possible to regenerate whole plants from the cultured tissue. On the other hand, mutagenised pollen may be applied to any recipient maize genotype, including commercially important and well-established elite breeding lines. 30 Also the rate of occurrence of undesirable mutations which somaclonal variation is known to produce is reduced.

In this invention, selection is carried out at at the M_1 generation, with the result that only

dominant mutations are selected. Also, being carried out on whole plants or on the seed pre-emergence, or both, allows the herbicide concentration to mimic the field conditions more closely than is possible with
5 the application of the selection pressure of the herbicide to a tissue culture. In the tissue culture selection method, whole plants have to be regenerated from the tissue and grown to maturity before any indication of the performance of the progeny under
10 field application rates of the herbicide can be obtained. In our method, selection is made directly on the plants under concentrations of herbicide which are comparable to those which are recommended for normal weed-killing activity in the field. Selection
15 on the M_2 generation, as with the tissue culture method, selects recessive mutations as well as dominant traits. Dominance of a desirable trait is generally viewed as more useful and easier to handle in a breeding programme especially of hybrid crops.

20 We believe that the mutants selected under herbicide pressure vary according to the particular member of the herbicide family which is used. All of our mutants were selected under pressure of imazethapyr. Had a different imidazolinone herbicide
25 been used, a different spectrum of mutants would have been selected.

We have found, quite surprisingly, that the mutants which we have isolated by the method of the invention are free of deleterious mutations. This was
30 entirely unexpected and the reason for this advantage is not entirely clear. We believe, but do not wish to be bound by this explanation, that the degree of control which we are able to exercise over the selection step, using carefully controlled rates of

application of the herbicide, for example, giving a good overall and uniform rate of exposure, may be responsible.

5 The invention will now be described by the following summary of the method by which the herbicide-resistant plants of the invention were derived.

The Figures which accompany this application are as follows:

10 Figure 1 is a flow-chart showing the derivation of several generations of progeny from plants generated by this invention;

Figure 2 shows the chemical structures of the herbicides used in this invention;

15 Figure 3 is a graph of ALS enzyme activity in the presence of imazethapyr. The enzyme is extracted from plants heterozygous for the resistance gene;

20 Figure 4 is a graph of ALS enzyme activity in the presence of imazapyr. The enzyme is extracted from plants heterozygous for the resistance gene;

Figure 5 is a graph of ALS enzyme activity in the presence of imazaquin. The enzyme is extracted from plants heterozygous for the resistance gene;

25 Figure 6 is a graph of ALS enzyme activity in the presence of chlorsulfuron. The enzyme is extracted from plants heterozygous for the resistance gene;

Figure 7 is a graph of ALS enzyme activity in the presence of chlorimuron. The enzyme is extracted from plants heterozygous for the resistance gene;

30 Figure 8 is a graph of ALS enzyme activity in the presence of thiocarburon. The enzyme is extracted from plants heterozygous for the resistance gene;

Figure 9 is an enzyme activity graph of the activity of ALS extracted from leaves of progeny of

mutants 1 and 2 which are homozygous for the resistant allele and in the presence of imazethapyr (PURSUIT);

Figure 10 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of imazaquin (SCEPTER);

Figure 11 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of imazapyr (ARSENAL);

Figure 12 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of chlorsulfuron (GLEAN);

Figure 13 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of chlorimuron (CLASSIC);

Figure 14 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of thiacarburon (HARMONY);

Figure 15 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of a triazolopyrimidine;

Figure 16 is an enzyme activity graph of the activity of ALS extracted from leaves of mutants 1 and 2 which are homozygous for the resistant allele and in the presence of a phenoxyprymidine;

Figure 17 is a dose response curve for the herbicide imazethapyr (PURSUIT);

Figure 18 is a dose response curve for the herbicide imazaquin (SCEPTER);

Figure 19 is a dose response curve for the herbicide chlorimuron (CLASSIC);

Figure 20 is a dose response curve for the herbicide chlorsulfuron (GLEAN);

5 Figure 21 is a dose response curve for a triazolopyrimidine herbicide;

Figure 22 is a dose response curve for a phenoxyypyrimidine herbicide; and,

10 Figure 23 is a dose response curve for the herbicide imazethapyr (PURSUIT) for mutants 1 and 2 in both heterozygous and homozygous forms.

1. PRODUCTION OF M1 SEED

A stock solution of ethyl methane sulphonate (EMS) was made up to contain one millilitre of EMS in 15 100 ml of light paraffin oil. The stock solution was stored under refrigeration.

20 Fresh pollen grains with anthers were harvested from a total of twenty tassels of field grown maize inbred line UE95. Pollen grains were separated from the anthers using a Glassine (Trade Mark) bag.

25 Around 3 milligrams of pollen were added to 45 millilitres of the EMS stock solution in a 60 ml capacity bottle. The pollen/EMS solution mixture was shaken vigorously for 30 seconds then shaken four or five times every three minutes over a period of 40 minutes, to prevent precipitation of the pollen grains. The treated pollen grains were brushed on to the silks of the detasseled inbred female parent (coded UE95).

30 The plants were grown to maturity and the M1 seeds harvested.

2. SCREENING

M1 seed was sown, 100 seeds per tray, in WCB growing medium (low organic matter, 45% loam, 55%

grit) and sprayed to 'run-off' using a track sprayer, with a solution of imazethapyr (PURSUIT) at a concentration calculated to be the equivalent of 250 g/ha of active ingredient. The seeds were covered 5 with 0.5 inch of WCB and grown in the glasshouse at 25°C.

The chosen application rate was such that germination was close to 100%, but subsequently all susceptible plants were severely affected. However, 10 after about two weeks, the initial effect of the herbicide became apparent: thin striped leaves and reduced height of only about 20% of the height of the control plants. After three to four weeks almost all the sensitive plants were completely dead. Unsprayed 15 UE95 plants were always grown in parallel with each screen as a control as it was already known that normal and M1 seedlings of UE95 are almost indistinguishable when germinated and grown to maturity without spraying.

20 3. SELECTION AND GROWTH

A total of ten plants (representing 0.01% of the total) were initially selected from the screen as exhibiting tolerance of normally lethal dosages of the herbicide. Mutant No.4 was subsequently found to be 25 of the sensitive phenotype and was withdrawn from the programme. Of the remaining nine, the majority were morphologically similar to the untreated control plants after selection but four were affected, being shorter and exhibiting other herbicidal effects. 30 Samples of seeds of these nine plants back-crossed to the parent UE95 (Mutants 1 to 10, excluding No.4) are those which have been deposited with the National Collection of Industrial and Marine Bacteria (see Table 1 above).

4. RFLP STUDY

In order to confirm that the sibling plants were indeed of UE95 genotype in origin and not an intrinsically more resistant contaminating inbred or 5 hybrid, RFLP (restriction fragment length polymorphism) fingerprinting analysis was performed on DNA extracted from leaf tissue from all ten plants.

Approximately 1 to 5 grams of leaf tissue was removed from each plant (ranging in age from four 10 weeks to 'mature'), DNA extracted and RFLP analysis performed using diagnostic single copy probes.

RFLP analysis confirmed that the selected plants were of the UE95 genotype.

5. SEGREGATION STUDY

15 Resistant plants were used in reciprocal back-crosses with homozygous, herbicide sensitive UE95. The frequency of resistant progeny in the M_1BC_1 or M_1BC_2 generations was found by treatment with imazethapyr and counting the survivors. The results 20 are shown in Table 2 below, along with the calculated value of χ^2 (for a 1:1 ratio).

As can be seen from the figures quoted in Table 2, the ratio of resistant to sensitive plants in the progeny of the backcross is not significantly 25 different from 1:1 for all of the mutants except numbers 3 and 7, indicating that resistance is controlled by a single dominant gene.

TABLE 2

Mutant No.	Number tested	Number sensitive	Number resistant	χ^2 (for 1:1)
1	40	18	22	0.40
2	228	110	118	0.28
3*	206	120	86	5.61
5	168	94	74	2.38
6	60	36	24	2.40
7*	168	139	29	72.00
8	188	88	100	0.77
9	168	83	85	0.02
10	134	73	71	0.03

* The resistance of these mutants to imazethapyr is low and it was therefore difficult to make a meaningful assessment in this experiment but, experience in other tests indicates that segregation is in the region of 1:1.

5 6. GENERATION OF F1 SEED FROM RESISTANT PLANTS

All ten plants identified in the screen were used in reciprocal sib-crossing with UE95 plants to give seed (designated M_1BC_1), that is, the resistant plants were used as both male and female donors.

10 7. RESCREENING OF THE PROGENY SEED

From each resulting cob, a small sample of seed was screened for imazethapyr (PURSUIT) resistance, employing the same conditions as are described above for the initial screen.

15 Resistant and sensitive phenotypes segregated in the progeny of each resistant plant.

8. PRODUCTION OF FURTHER GENERATIONS

Referring to Figure 1, the M_1BC_1 plants were self pollinated to give generation $M_1BC_1S_1$ which was again self-pollinated to give $M_1BC_1S_2$. From that generation 5 it was possible to identify, by the fact that the resistant trait is non-segregating, lines which are homozygous for the trait. These homozygous lines derived from Mutant No.1 may be utilised for the production of F1 hybrids which possess resistance to 10 the imidazolinone herbicides or for further breeding work.

9. ENZYME ASSAYS

Seeds of generation M_1BC_1 M_1BC_2 of the nine 15 mutants were sprayed pre-emergence with the herbicide imazethapyr (PURSUIT) (Trade Mark) at a rate of 250 g/ha and grown in a growth chamber (16 hour day, 27°C; 8 hour night, 17°C). Plants were harvested after 11 days.

Four grams of leaf material were harvested from 20 just above the first leaf axis and ground in a mortar and pestle in 20ml of a solution containing 40mM Tricine (Trade Mark), 10mM EDTA, 5mM pyruvate, 80 μ M flavin adenine dinucleotide (FAD), 1mM dithiothreitol (DTT), pH 8, plus 0.8g Polyclar AT. The homogenate 25 was pressed through four layers of muslin and centrifuged at 30,000g for 20 minutes.

The supernatant was brought to 65% saturation with ammonium sulphate, left to precipitate for 30 minutes and then centrifuged at 30,000g for 20 30 minutes. The pellet was resuspended in 2.5ml of 40mM Tricine, 10mM EDTA, 5mM pyruvate, 80 μ M FAD, 25% (v/v) glycerol, 1mM DTT, pH 8, and desalting into 3.5ml of 40mM Tricine, 10mM EDTA, 25% (v/v) glycerol, 1mM DTT, pH 8.

One hundred microlitres of the enzyme extract was used for each assay. Final reagent concentrations were: 120mM Tricine, 50mM pyruvate, 10mM $MgCl_2$, 66mM FAD, 93 μ M thiamine pyrophosphate (TPP), pH 8, in the presence or absence of a herbicide of interest, in a volume of 750 μ l. Incubations were carried out at 37°C for 30 minutes. Reactions were stopped with 250 μ l of 1.84M sulphuric acid, and decarboxylation of the acetolactate carried out at 37°C for 75 minutes, or at 60°C for 30 minutes. Assay blanks had sulphuric acid added prior to the enzyme extract. To the samples and blanks 650 μ l of α -naphthol in 2.73M NaOH/0.16% (w/v) creatine were added followed by incubation at 37°C for 30 minutes. After centrifugation at 30,000g the optical density of the supernatant was read at 540nm.

The assay procedure described was carried out on ALS enzyme extracted from each of the nine mutants for the following herbicides:

- 20 imazethapyr (PURSUIT)
- imazapyr (ARSENAL)
- imazaquin (SCEPTER)
- chlorsulfuron (GLEAN)
- chlorimuron (CLASSIC)
- 25 thiocarburon (HARMONY)

The enzyme activity graphs are given as Figures 3 to 8. In addition, homozygous lines of mutants 1 and 2 which were selected from generation $M_1BC_1S_2$, were also tested against the same group of herbicides and 30 against representatives of the triazolopyrimidines and phenoxyprymidines. The results are shown in Figures 9 to 16. The chemical structures of these herbicides are shown in Figure 2 of the accompanying drawings.

10. ASSESSMENT OF INHIBITION

From the enzyme assay data it is possible to derive, as a measure of the degree of inhibition the factor ID_{50%} for each mutant and for each 5 herbicide. This factor is indicative of the herbicide concentration which gives 50% inhibition of the ALS activity in comparison with the wild-type control. It is also possible to derive the fold increase in resistance. These 10 calculations are given in Table 3 below.

TABLE 3

IMAZETHAPYR (heterozygous mutants)

Plant	ID50% μM	Fold Increase
Wild type	1.5	
Mutant 1	>1000	>660
2	230	153
3	2.5	2
5	125	83
6	60	40
7	4	3
8	>1000	>660
9	425	283
10	15	10

IMAZAPYR (heterozygous mutants)

Plant	ID50% μM	Fold Increase
Wild type	3	
Mutant 1	>1000	>333
2	540	180
3	4	1
5	300	100
6	5-	2
7	10	3
8	>1000	>333
9	>1000	>333
10	35	12

IMAZAQUIN (heterozygous mutants)

Plant	ID50% μM	Fold Increase
Wild type	0.6	
Mutant 1	150	250
2	20	33
3	1.5	2
5	25	42
6	1.5	2
7	1	2
8	150	250
9	50	83
10	7	12

CHLORIMURON (heterozygous mutants)

Plant	ID50% nM	Fold Increase
Wild type	3.7	
Mutant 1	5	1
2	10	3
3	15	4
5	17	5
6	15	4
7	5	1
8	10	3
9	10	3
10	17	5

CHLORSULFURON (heterozygous mutants)

Plant	ID50% nM	Fold Increase
Wild type	18.5	
Mutant 1	65	4
2	65	4
3	35	2
5	40	2
6	65	4
7	8	1
8	75	4
9	50	3
10	35	2

THIACARBURON (heterozygous mutants)

Plant	ID50% nM	Fold Increase
Wild type	42.5	
Mutant 1	55	1
2	320	8
3	85	2
5	340	8
6	45	1
7	25	1
8	40	1
9	45	1
10	390	9

Similar calculations were made for homozygous mutants 1 and 2 only, the results being given in Table 4 below.

TABLE 4

MUTANT 1 (homozygous)

Herbicide	ID50%		Fold Increase
	Wild	Mutant	
Imazapyr	3 μ M	>1mM	>333
Imazethapyr	1.5 μ M	750 μ M	500
Imazaquin	0.6 μ M	175 μ M	292
Chlorimuron	3.7nM	10nM	2.7
Chlorsulfuron	18.5nM	60nM	3.2
Thiacarburon	42.5nM	40nM	nil

TABLE 4 (continued)

MUTANT 2 (homozygous)

Herbicide	Wild	ID50% Mutant	Fold Increase
Imazapyr	3 μ M	675 μ M	225
Imazethapyr	1.5 μ M	400 μ M	267
Imazaquin	0.6 μ M	40 μ M	66
Chlorimuron	3.7 nM	20 nM	5.4
Chlorsulfuron	18.5 nM	40 nM	2.2
Thiacarburon	42.5 nM	360 nM	8.5
Triazolo- pyrimidine	0.07 μ M	0.12 μ M	1.7
Phenoxy- pyrimidine	0.03 mM	0.45 mM	22.5

11. CROSS-RESISTANCE TESTING

5 The seeds used in this screen were populations of mixed tolerant and sensitive seed, segregating 1:1, derived from each of the heterozygous, tolerant mutants crossed with homozygous, sensitive UE95. All of the tolerant progeny are heterozygous for tolerance. Each of the mutants identified above were used except mutant 4. The controls were seed of self-pollinated UE95 which was pollinated 10 in the same year as the tolerant lines.

15 One litre of compost was placed in each seed tray of dimensions 19cm x 11cm x 5cm deep and firmed down flat. Two furrows 1 cm deep were drawn in the surface of the compost in each tray and six seeds were sown in each furrow (total twelve seeds per tray). Some trays were sown with 24 seeds, in which case three furrows were made and 8 seeds sown per furrow.

In each test each tray contained either the seed of one mutant or of the UE95 control for each herbicide application rate and for the untreated control.

5 Five different types of ALS-inhibitor herbicides were tested on the mutants plus imazethapyr (PURSUIT) as a comparison. Four rates of each were applied; approximately 0.1x, 0.5x, 1x and 4x the estimated field rates for each 10 herbicide. The rates are the same for mutant 3 to 10 but higher rates for chlorimuron (CLASSIC) and lower rates for chlorsulfuron (GLEAN) were applied to mutants 1 and 2.

15 The following herbicides, with their estimated field application rates were used:

chlorimuron (CLASSIC)	8 -13 g/Ha
chlorsulfuron (GLEAN)	4 -26 g/HA
imazaquin (SCEPTER)	100 -150 g/Ha
triazolopyrimidine	10 - 30 g/Ha
20 phenoxypyrimidine	100 - 200 g/Ha
imazethapyr (PURSUIT)	70 - 140 g/Ha

The rates of application used in the screening tests were as follows:

25 chlorimuron (CLASSIC) 4,20,40,160 g/Ha (mutants 1- and 2); and, 8,40,80,320 g/Ha (mutants 3 to 10);
chlorsulfuron (GLEAN) 5,25,50,200 g/Ha (mutants 1 and 2); and, 1,5,10,40 g/Ha (mutants 3 to 10);
imazaquin (SCEPTER) 30,150,300,1200 g/Ha;
triazolopyrimidine 5,25,50,200 g/Ha;
30 phenoxypyrimidine 10,50,100,400 g/Ha; and,
imazethapyr (PURSUIT) 30,150,300,1200 g/Ha

5 The triazolopyrimidine and phenoxyprymidine were formulated in an adjuvant wetting agent known as JF5969 and diluted with water to make a final concentration of 10% JF5969. The remaining compounds were diluted with water only.

10 The spray jet and parameters were as described above under "SCREENING". After spraying, the seeds were covered with 0.5 litre of compost and firmed down flat (2.5cm covering) and grown under the conditions described under "SCREENING" above.

15 Visual assessments were made of the plants after four weeks growth. Each plant was scored on a scale of zero to 5 and the height measured from soil surface to the tallest leaf tip. The scoring scale was as follows:

0 - 0 to 10% damage (little or no herbicidal effect)
1 - 11 to 25% damage
20 2 - 26 to 50% damage
3 - 51 to 85% damage
4 - 81 to 95% damage
5 - 96 to 100% damage (complete death of the plant)

25 In carrying out these assessments, plants which were obviously of the sensitive phenotype were ignored (approximately 50% of the plants).

Plants scoring 0, 1 or 2 were recorded and potted on into larger pots and grown to maturity.

30 The results are shown in Figures 17 to 22. Figure 23 shows a comparison of the results obtained as between heterozygous and homozygous plants.

12. INTERPRETATION OF THE RESULTS

The data generated for the heterozygotes by the enzyme assays and the dose response curves on the living plants, do not correlate precisely. To those skilled in the art, this will not be particularly unexpected. However, there is a general correlation in the sense that the general level of resistance shown in enzyme assays tends to be reflected in the glasshouse studies. Of course, the criteria for deciding what constitutes a useful mutant varies according to the herbicide, or spectrum of herbicides and the rate of application of the herbicide with which one is dealing and the relatively rich diversity in the spectrum of cross-resistance displayed by the mutants of this invention is seen as an advantage rather than an undesirable variation, allowing, as it does, selection of a mutant for the circumstances of intended use. For example, although a particular mutant may display relatively low resistance compared with the others against a particular herbicide this may make it eminently useful for providing plants which are intended to tolerate only small amounts of herbicide, for example, to resist a "carry-over" effect.

With this in mind, it is therefore possible to categorise the mutants on the basis of the enzyme assays and the glasshouse tests into "strong", "intermediate", "weak" and "zero" resistance groups for each herbicide. This classification is summarised in Table 5 below.

TABLE 5

IMAZETHAPYR		MUTANT NUMBER	
Strong	Enzyme Assay		Dose Response
Intermediate	1,8,9		1,5,8,9
Weak	2,5,10		2,6,10
Zero	3,6,7,		7,3
	-		-
IMAZAPYR		Enzyme Assay	Dose Response
Strong	1,2,5,8,9,10		no data
Intermediate	6,7		no data
Weak	3		no data
Zero	-		no data
IMAZAQUIN		Enzyme Assay	Dose Response
Strong	1,8,9		1,2,8,9
Intermediate	2,5		5
Weak	3,6,7,10		3,6,10
Zero			7
CHLORSULFURON		Enzyme Assay	Dose Response
Strong	-		-
Intermediate	-		3
Weak	1,2,3,5,6,8,9,10		5,7,9,10
Zero	7		1,2,6,8,9,10
CHLORIMURON		Enzyme Assay	Dose Response
Strong	-		-
Intermediate	10		-
Weak	1,2,3,5,6,7,9		3,10
Zero	8		1,2,5,6,7,8,9
THIACARBURON		Enzyme Assay	Dose Response
Strong	-		no data
Intermediate	2,5,10		no data
Weak	1,3,6,8,9		no data
Zero	7		no data

TABLE 5 (continued)

TRIAZOLOPYRIMIDINE		
Strong	Enzyme Assay	Dose Response
Intermediate	no data	3
Weak	no data	1,2
Zero	no data	7,8,9
		5,6,10
PHENOXYPYRIMIDINE		
Strong	Enzyme Assay	Dose Response
Intermediate	no data	-
Weak	no data	2,5,7
	no data	3,8,9,10
	no data	1,6

13. PLANT BREEDING

The mutanted lines of the present invention can be used in common with various second parent lines to produce herbicide resistant hybrids.

5 Material from the homozygous lines may be entered into a breeding programme involving further outcrossing, selfing, visual selection and herbicide screening in order to produce a range of new herbicide-tolerant hybrid seed.

10 The herbicide resistance trait can be transferred to new lines by the described mutation breeding approach or by conventional breeding practices, using selection for herbicide resistance as described hereinabove. Biochemical

15 and molecular screening techniques are also available to those skilled in the art to aid the process. The use of genetic engineering techniques are readily conceivable to isolate and transfer the resistance gene.

20 The objective of a breeding programme may simply be the beneficial transfer of herbicide resistance or may be more complex, involving concurrent improvement of agronomic performance.

CLAIMS:

1. A method of producing mutant Zea mays possessing resistance to a normally lethal concentration of a chosen herbicide, comprising subjecting isolated pollen of Zea mays to the action of a chemical mutagen, recovering mutagenised pollen, pollinating a female parent therewith, harvesting seed from the mature plant, growing the seeds in the presence of selection pressure of the chosen herbicide or a second herbicide having the same mode of action as the chosen herbicide and recovering plants and their progeny which are resistant to the applied selection pressure.
2. A method as claimed in claim 1 in which the chemical mutagen is ethyl methanesulphonate.
3. A method as claimed in claim 1 in which the herbicide utilised for applying selection pressure is an inhibitor of acetolactate synthase.
4. A method as claimed in claim 3 in which the said herbicide is imazethapyr.

5. A method as claimed in any preceding claims, in which the concentration of herbicide used for selection pressure is insufficient to inhibit germination but sufficient to inhibit growth of sensitive plants.

6. A method as claimed in any preceding claim in which the selection pressure is applied by application of the herbicide to the seeds pre-emergence.

7. Herbicide resistant Zea mays seed, samples of which has been deposited, under the terms of the Budapest Treaty, at the National Collection of Industrial & Marine Bacteria Ltd. Aberdeen, United Kingdom under the following Accession Numbers and on the dates indicated:

<u>Accession Number</u>	<u>Date of Deposit</u>
NCIMB 40137	8th May 1989
NCIMB 40136	8th May 1989
10 NCIMB 40138	8th May 1989
NCIMB 40167	18th July 1989
NCIMB 40168	18th July 1989
NCIMB 40169	18th July 1989
NCIMB 40170	18th July 1989
15 NCIMB 40171	18th July 1989
NCIMB 40172	18th July 1989

8. Imazethapyr resistant Zea mays seed which has been deposited, under the terms of the

5

Budapest Treaty, at the National Collection of Industrial & Marine Bacteria Ltd. Aberdeen, United Kingdom under the following Accession Numbers and on the dates indicated:

10

<u>Accession Number</u>	<u>Date of Deposit</u>
NCIMB 40137	8th May 1989
NCIMB 40167	18th July 1989
NCIMB 40170	18th July 1989
NCIMB 40171	18th July 1989

5

9. Imazaquin resistant Zea mays seed which has been deposited, under the terms of the Budapest Treaty, at the National Collection of Industrial & Marine Bacteria Ltd. Aberdeen, United Kingdom under the following Accession Numbers and on the dates indicated:

10

<u>Accession Number</u>	<u>Date of Deposit</u>
NCIMB 40137	8th May 1989
NCIMB 40136	8th May 1989
NCIMB 40170	18th July 1989
NCIMB 40171	18th July 1989

5

10. Imazapyr resistant Zea mays seed which has been deposited, under the terms of the Budapest Treaty, at the National Collection of Industrial & Marine Bacteria Ltd. Aberdeen, United Kingdom under the following Accession Numbers and on the dates indicated:

10

<u>Accession Number</u>	<u>Date of Deposit</u>
NCIMB 40137	8th May 1989
NCIMB 40136	8th May 1989
NCIMB 40167	18th July 1989
NCIMB 40170	18th July 1989
NCIMB 40171	18th July 1989

11. Herbicide resistant Zea mays plants derived from herbicide resistant seeds claimed in any of claims 7 to 10.
12. Herbicide resistant Zea mays hybrids created from the plants claimed in claim 11.
13. An allele of acetolactate synthase giving herbicide resistance which has been isolated from any one of the seeds claimed in claim 7.
14. An allele of acetolactate synthase giving herbicide resistance which is identical to an allele present in any one of the seeds claimed in claim 7.

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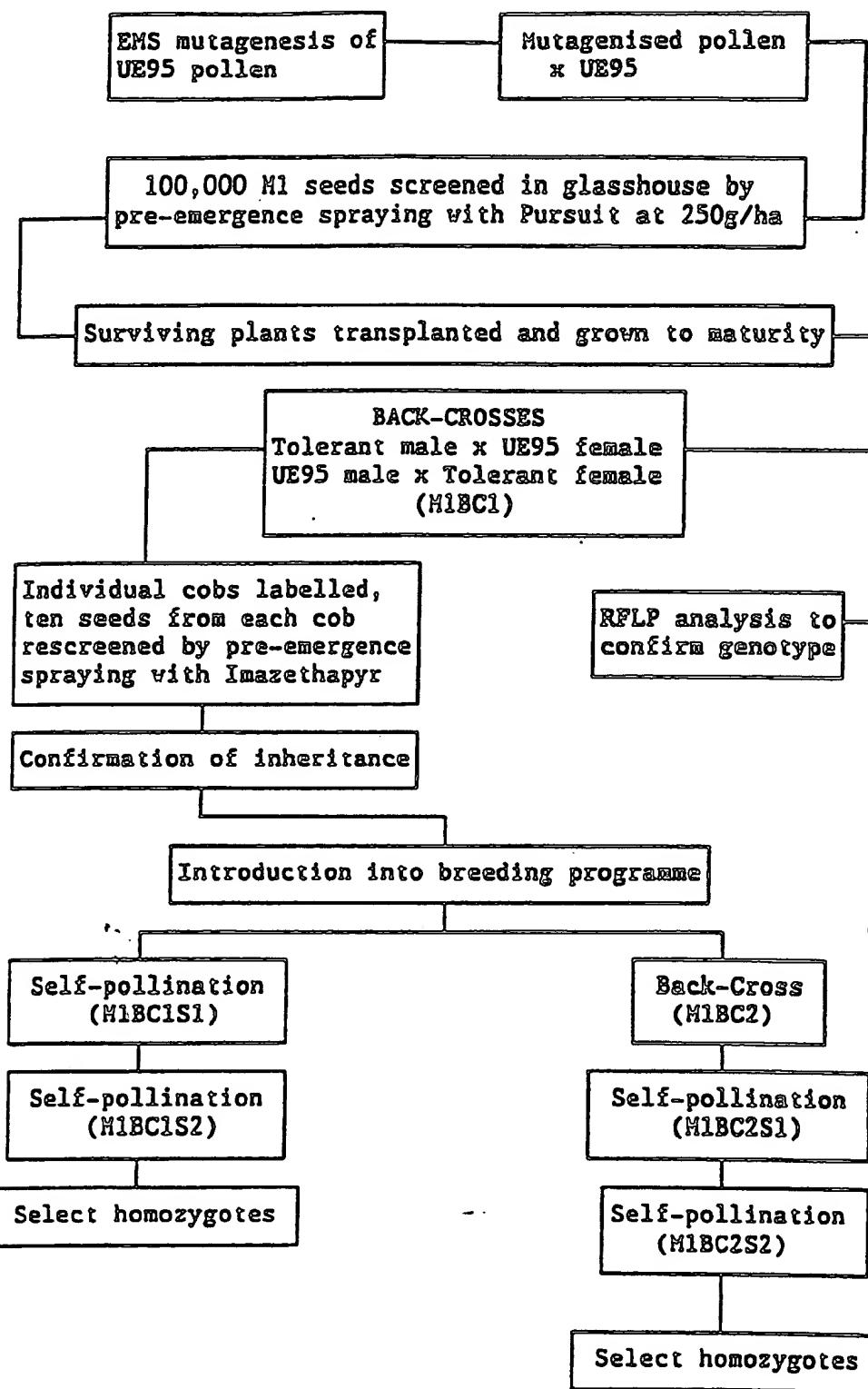
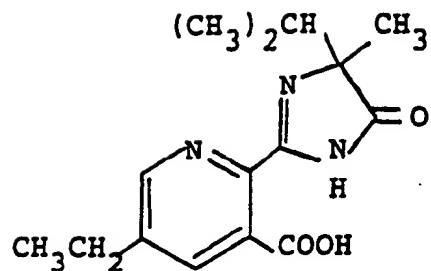
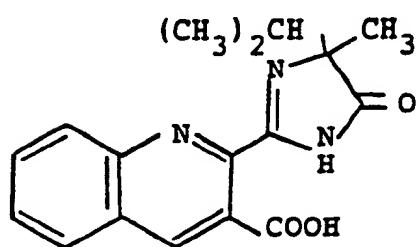


FIG. 1.

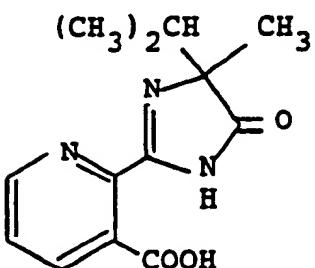
2/22



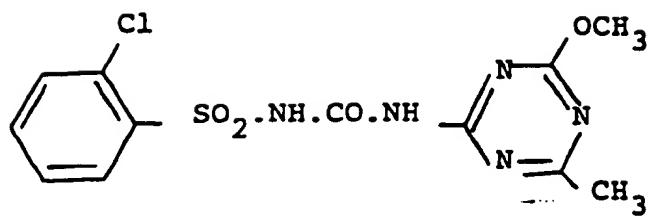
IMAZETHAPYR



IMAZAQUIN



IMAZAPYR



CHLORSULFURON

FIG. 2.

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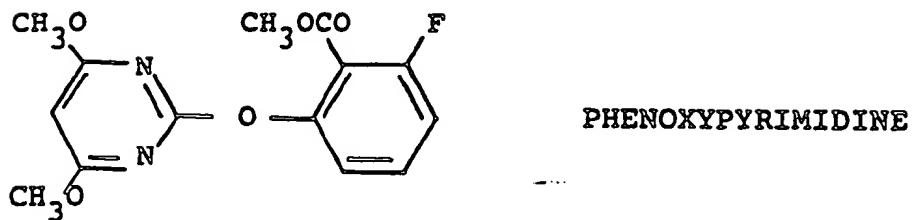
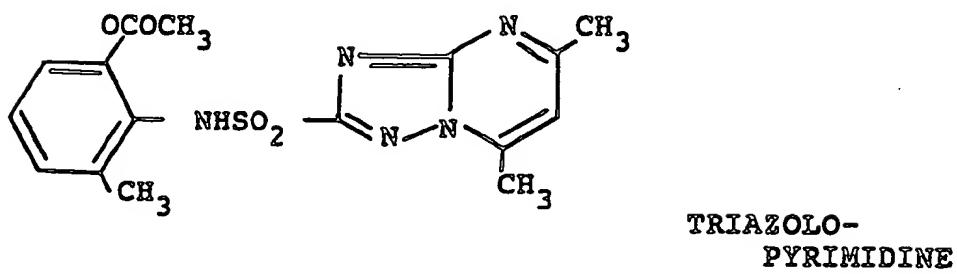
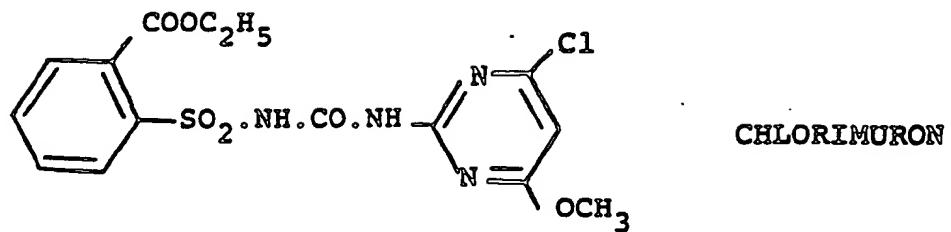
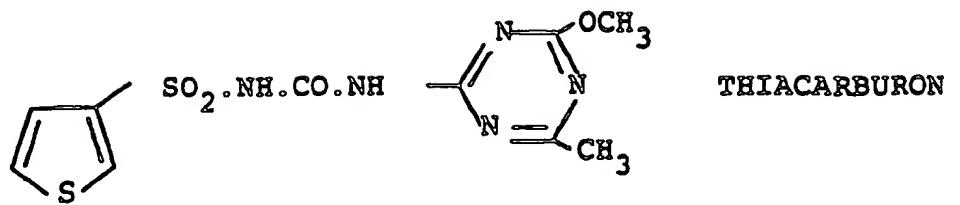


FIG. 2 Continued

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RESPONSE OF LEAF ALS TO 'PURSUIT'

HETEROZYGOTES

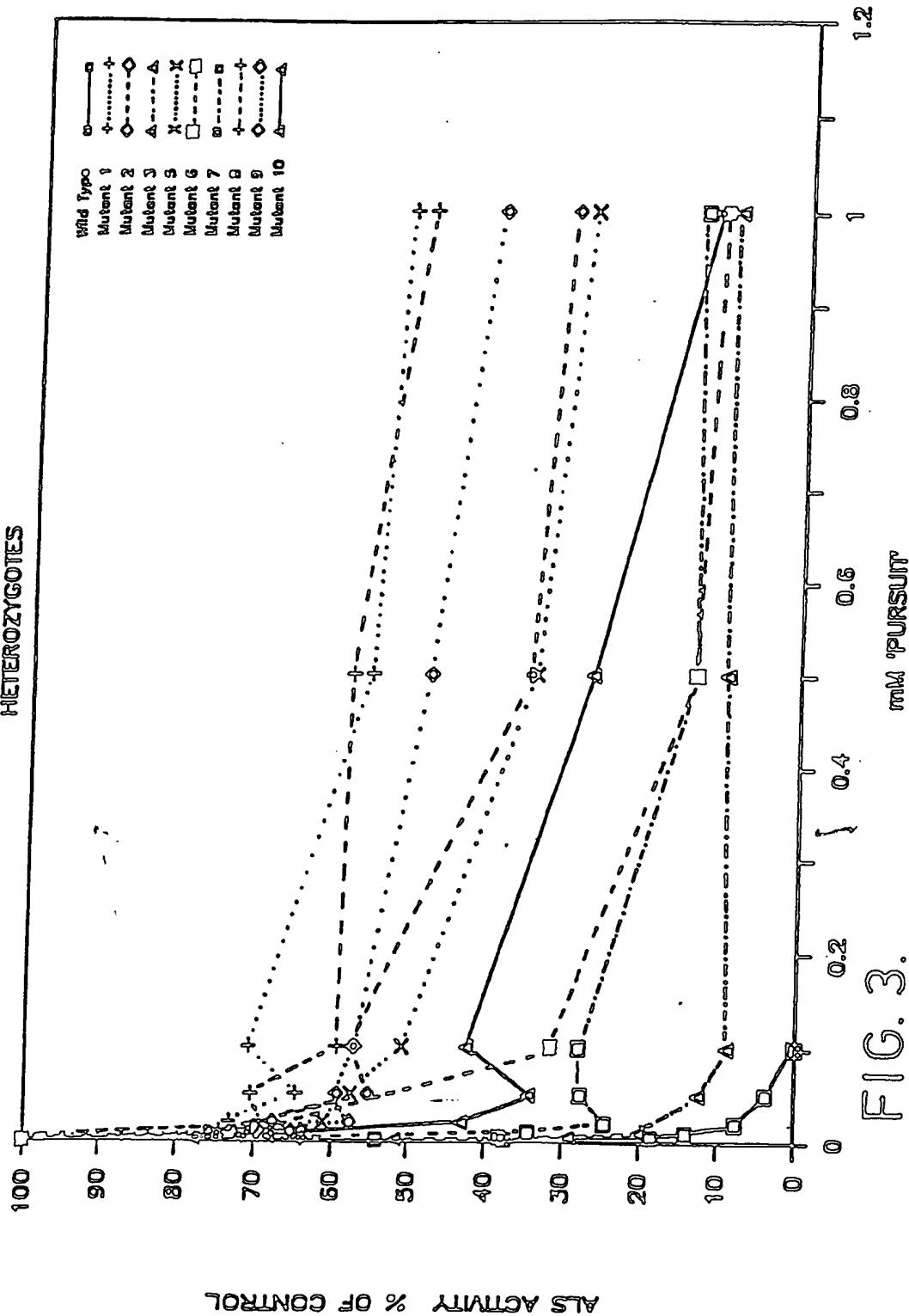


FIG. 3.

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RESPONSE OF LEAF ALS TO 'ARSENAL'¹

HETEROZYGOTES

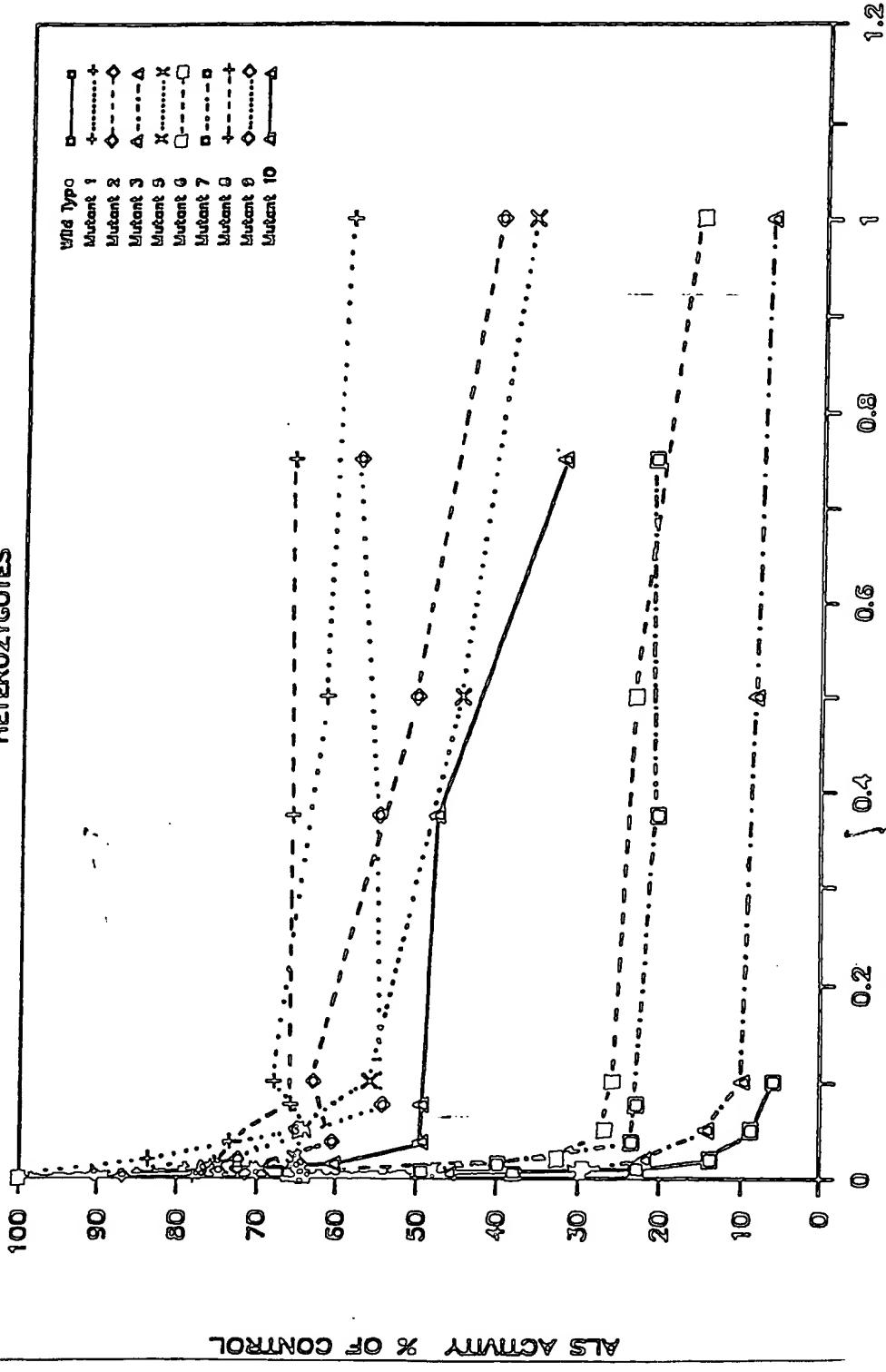


FIG. 4. MM 'ARSENAL'

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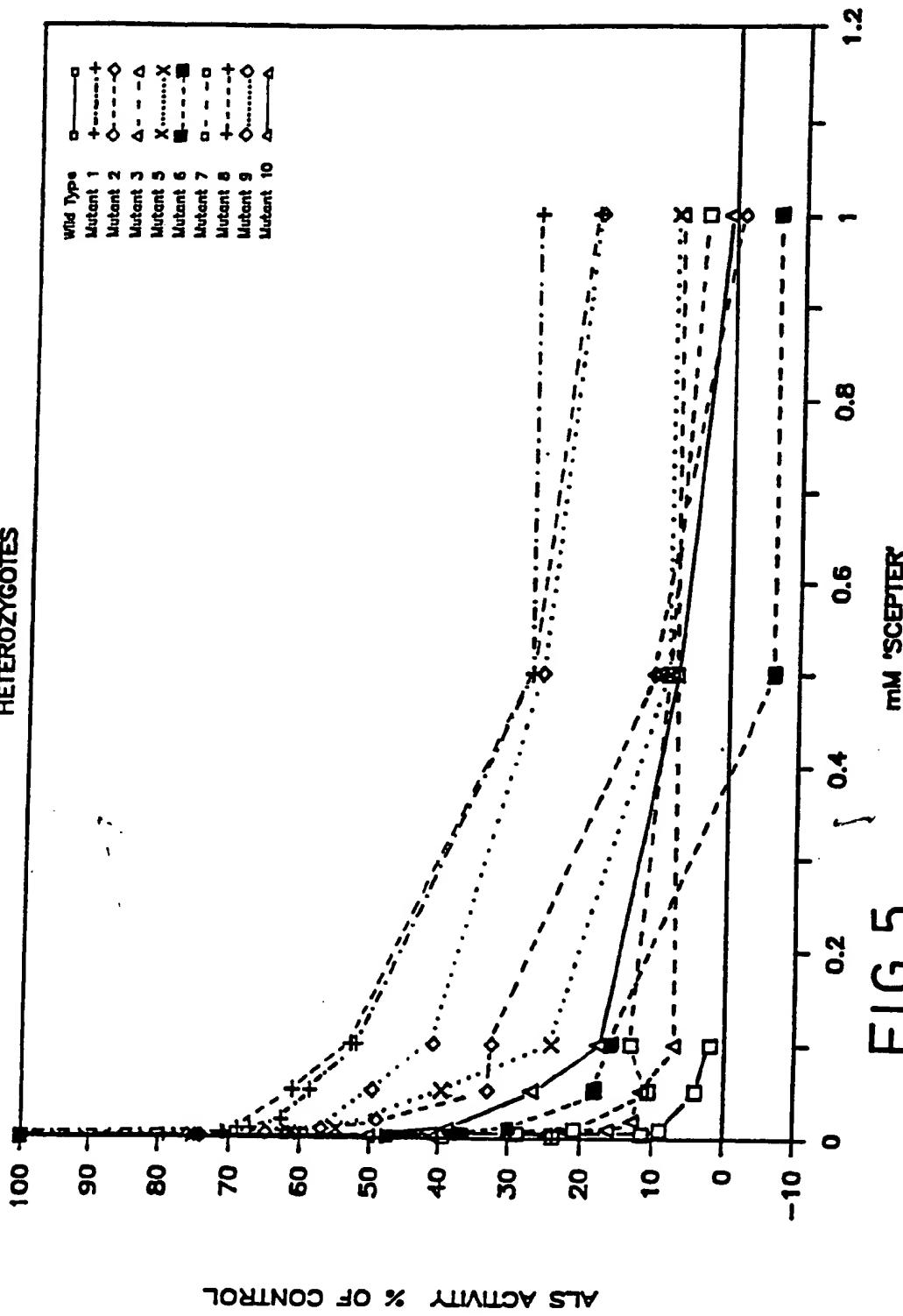
RESPONSE OF LEAF ALS TO 'SCEPTER'
HETEROZYGOTES

FIG. 5.

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RESPONSE OF LEAF ALS TO "GLEAN"

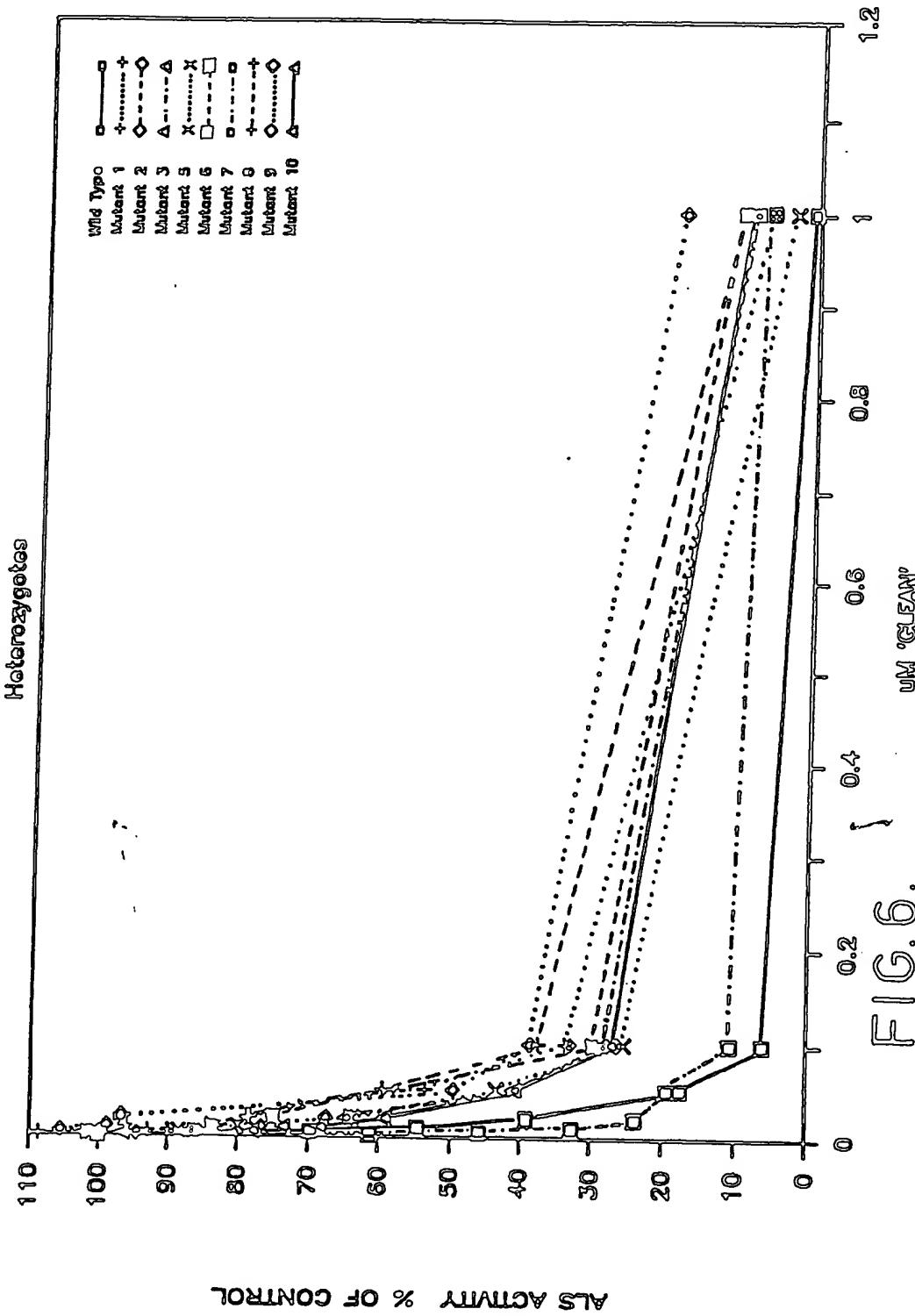


FIG. 6. uM "GLEAN"

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RESPONSE OF LEAF ALS TO 'CLASSIC'

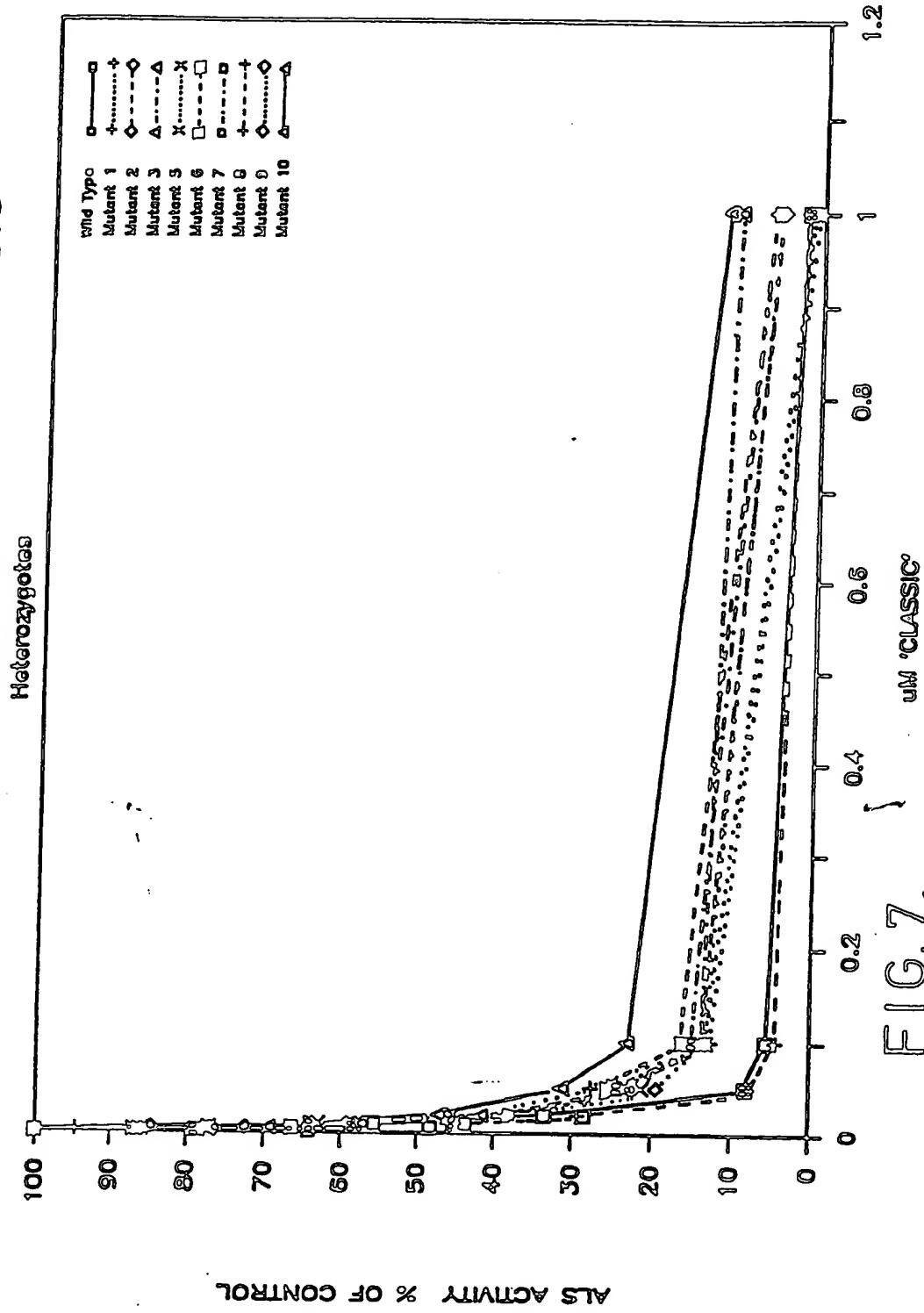


FIG. 7. 1 μM 'CLASSIC'

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RESPONSE OF LEAF ALS TO 'HARMONY'

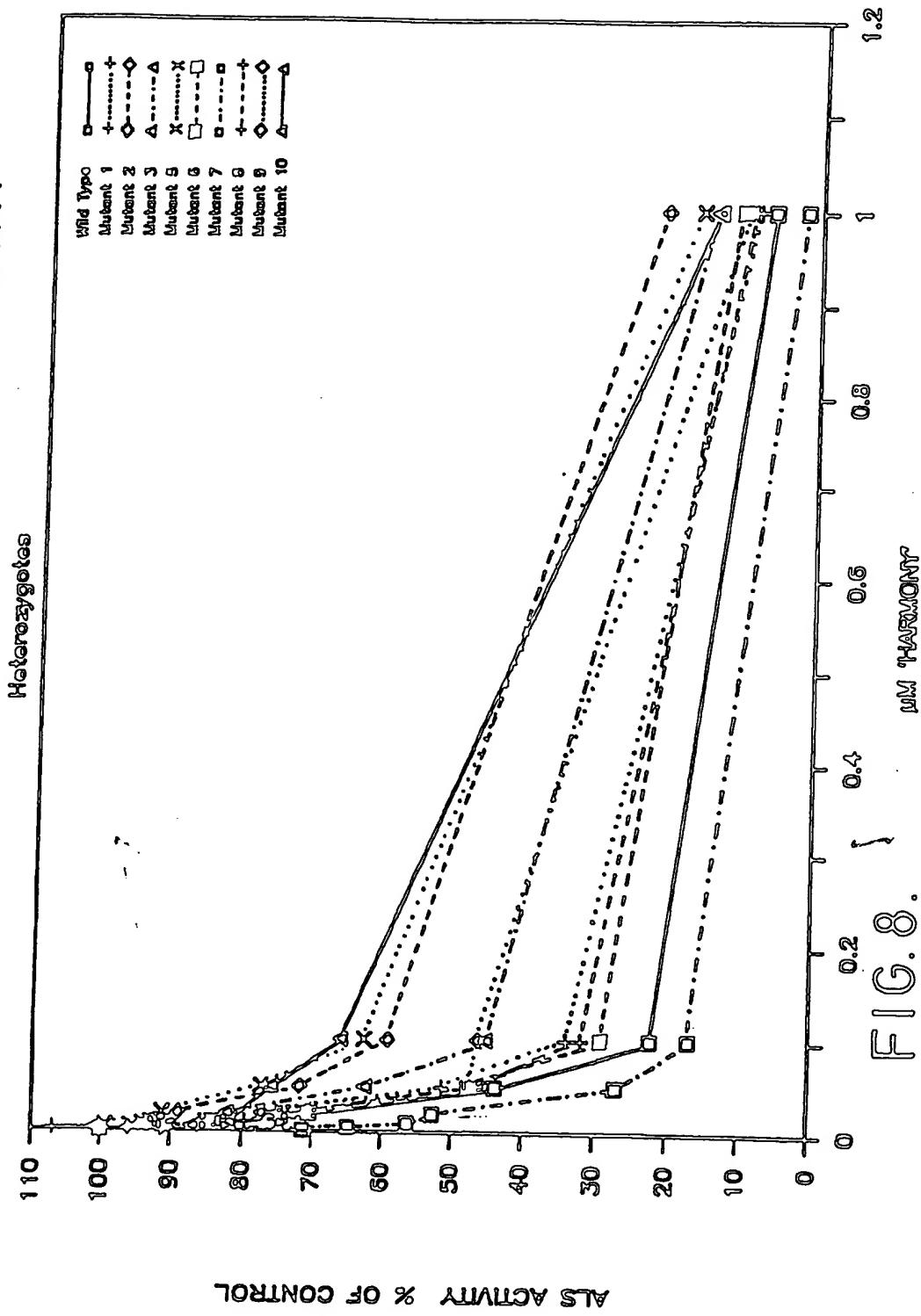


FIG. 8. MM 'HARMONY'

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RESPONSE OF LEAF ALS TO "PURSUIT"

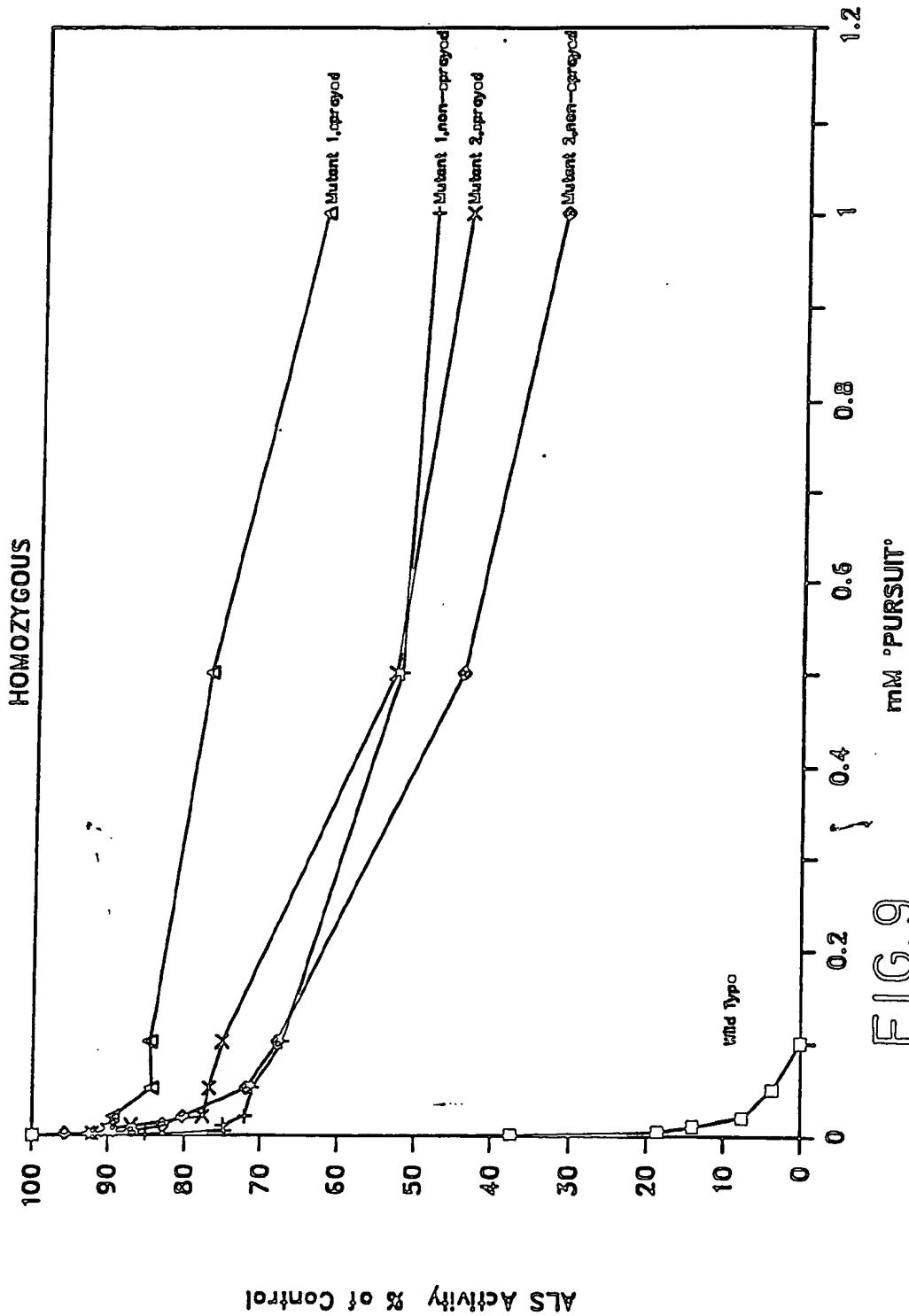


FIG. 9.

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RESPONSE OF LEAF ALS TO 'SCEPTER'

HOMOZYGOUS

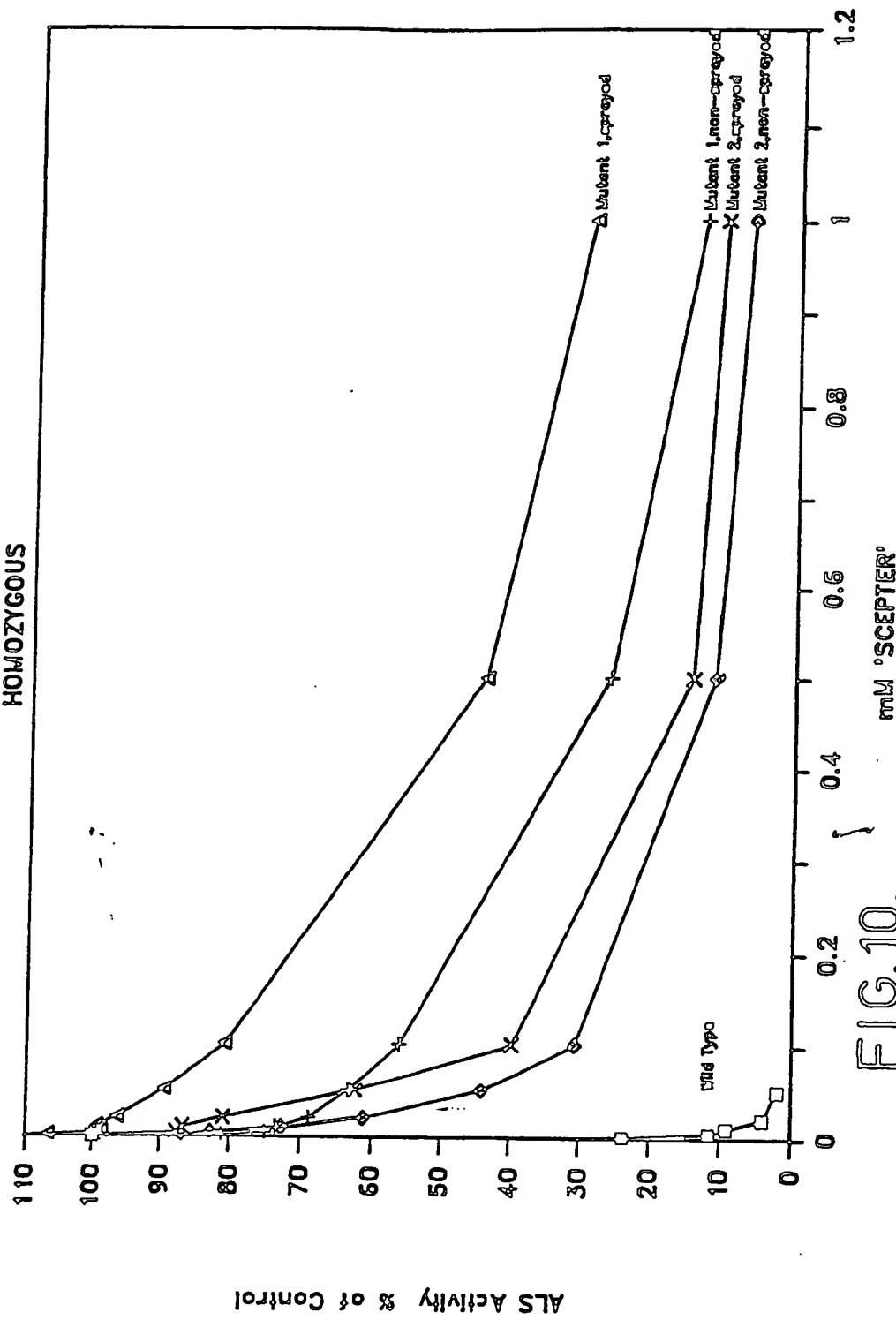


FIG. 10.

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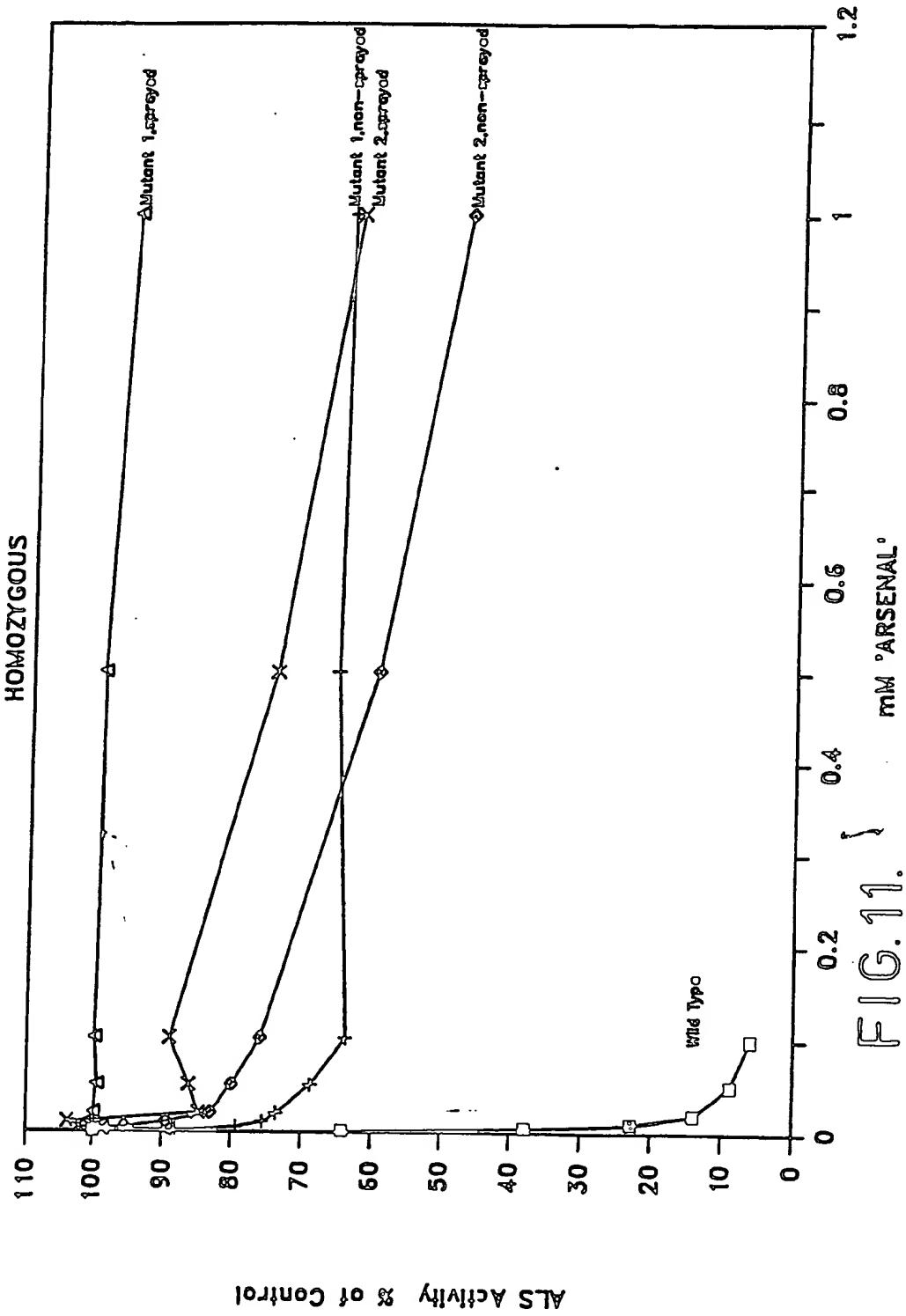
RESPONSE OF LEAF ALS TO 'ARSENAL'⁹

FIG. 11.

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FIG. 12. RESPONSE OF LEAF ALS TO 'GLEAN'

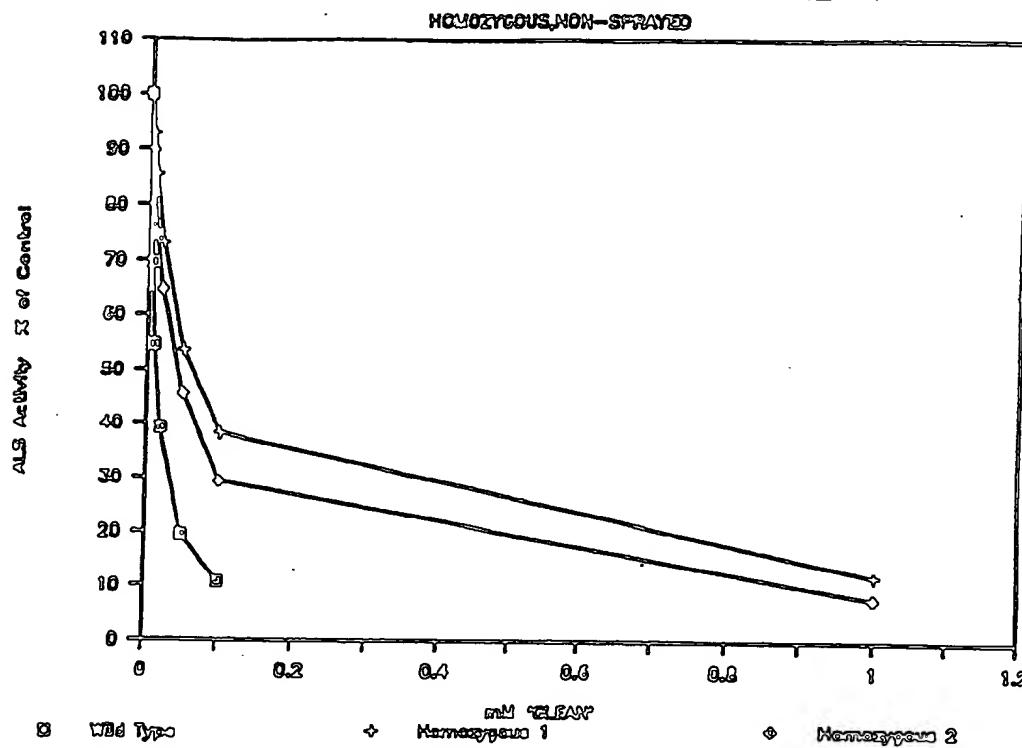
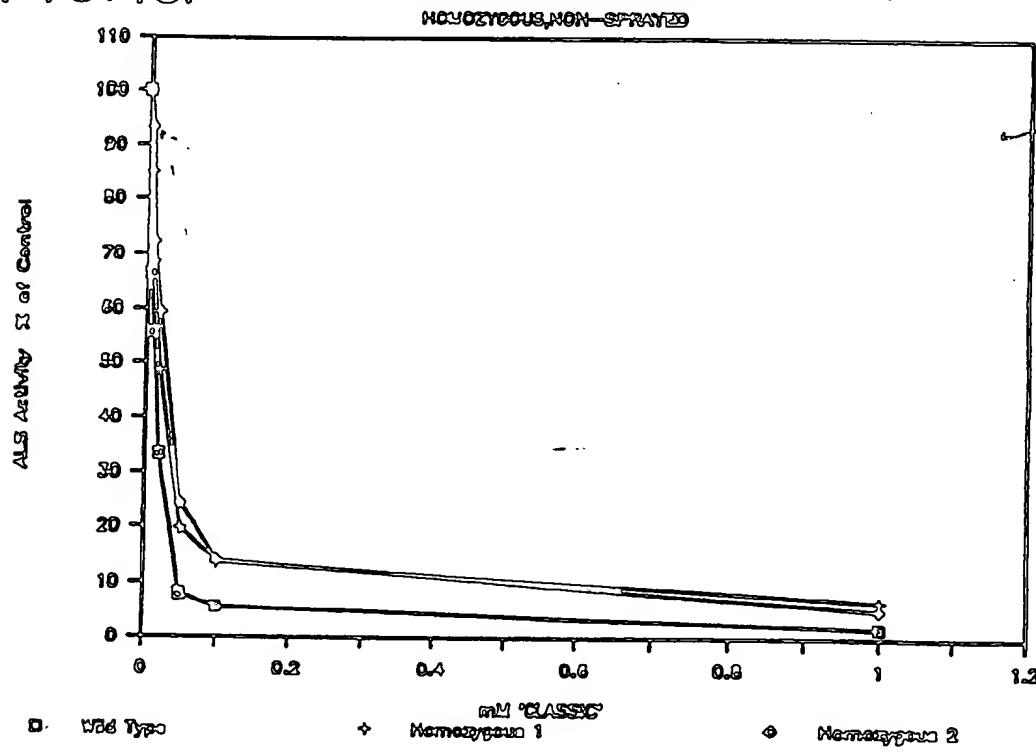
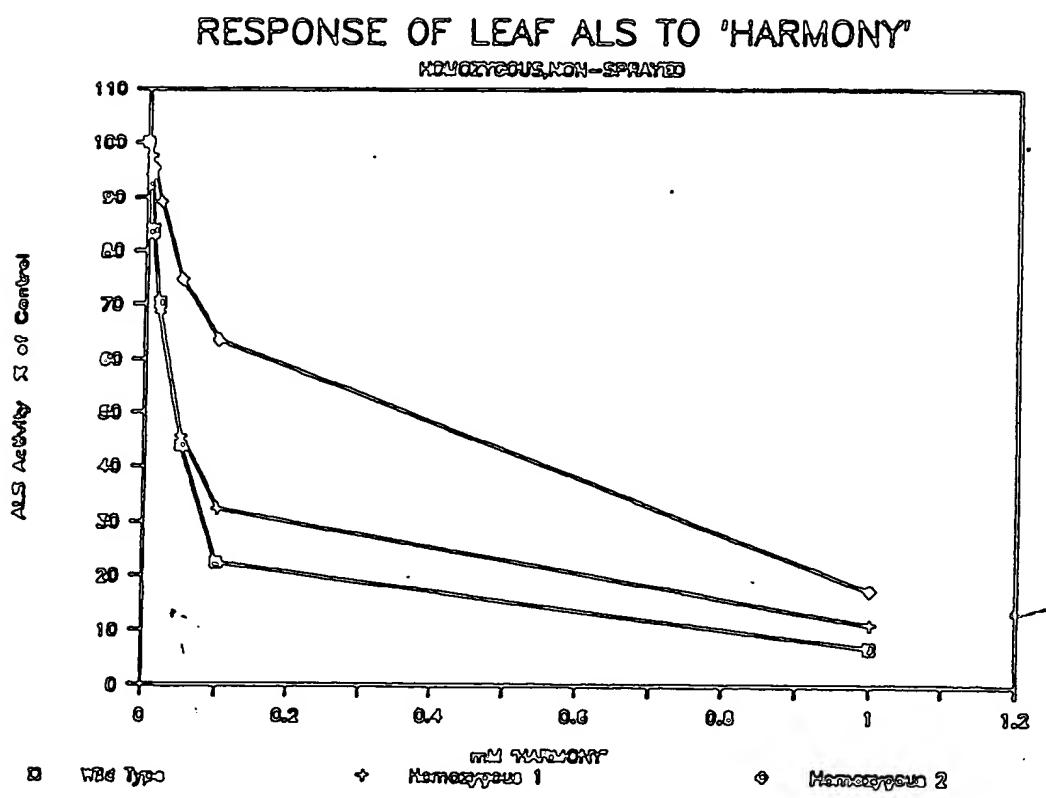


FIG. 13. RESPONSE OF LEAF ALS TO 'CLASSIC'



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FIG. 14.



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RESPONSE OF ALS TO A PHENOXPYRIMIDINE

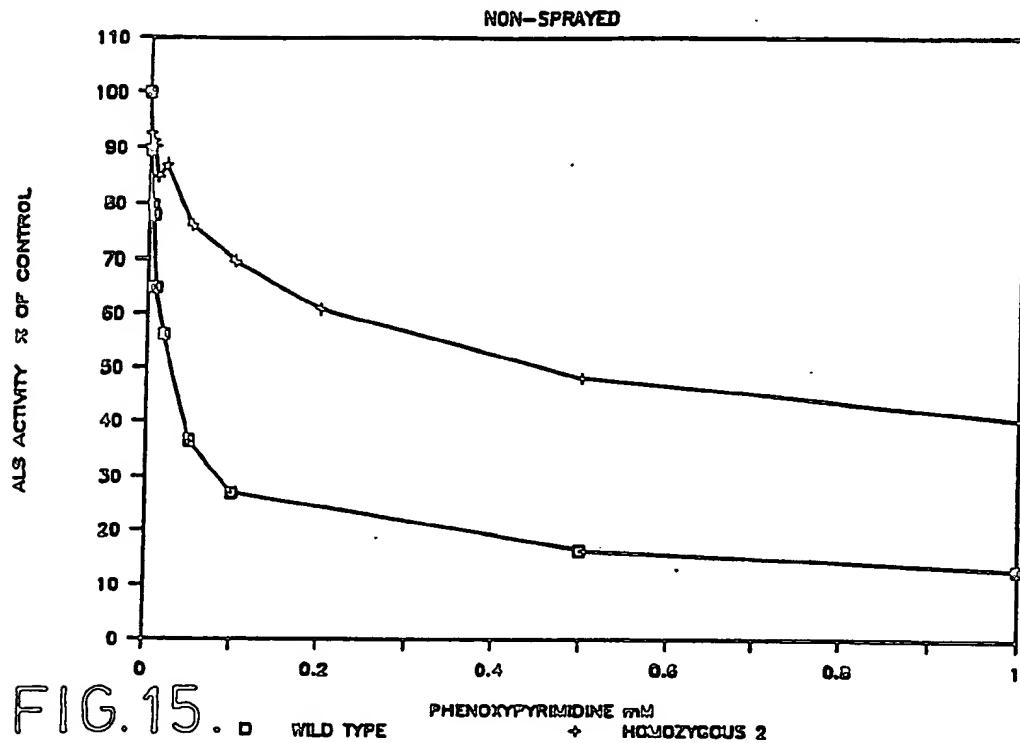


FIG. 15. □ WILD TYPE + HOMOZYGOUS 2

RESPONSE OF ALS TO A TRIAZOLOPYRIMIDINE

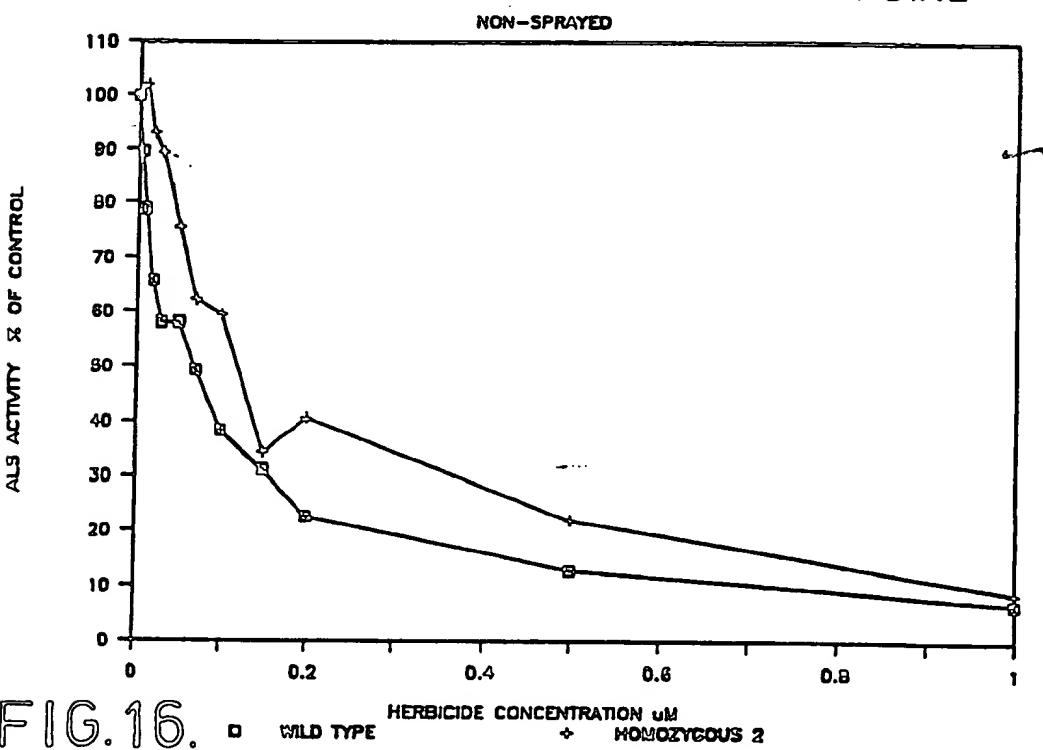
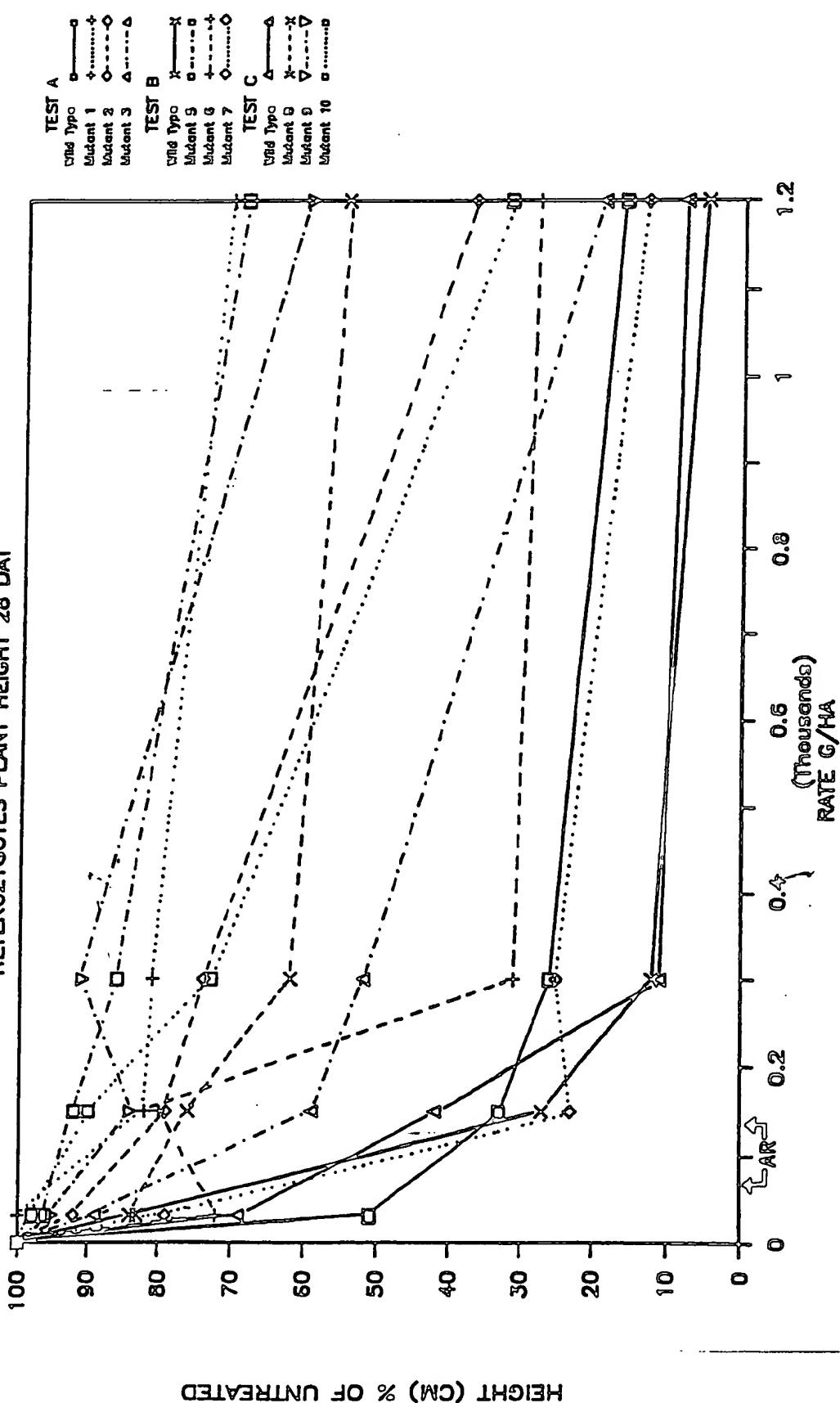


FIG. 16. □ WILD TYPE + HOMOZYGOUS 2

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FIG. 17. PURSUIT DOSE RESPONSE

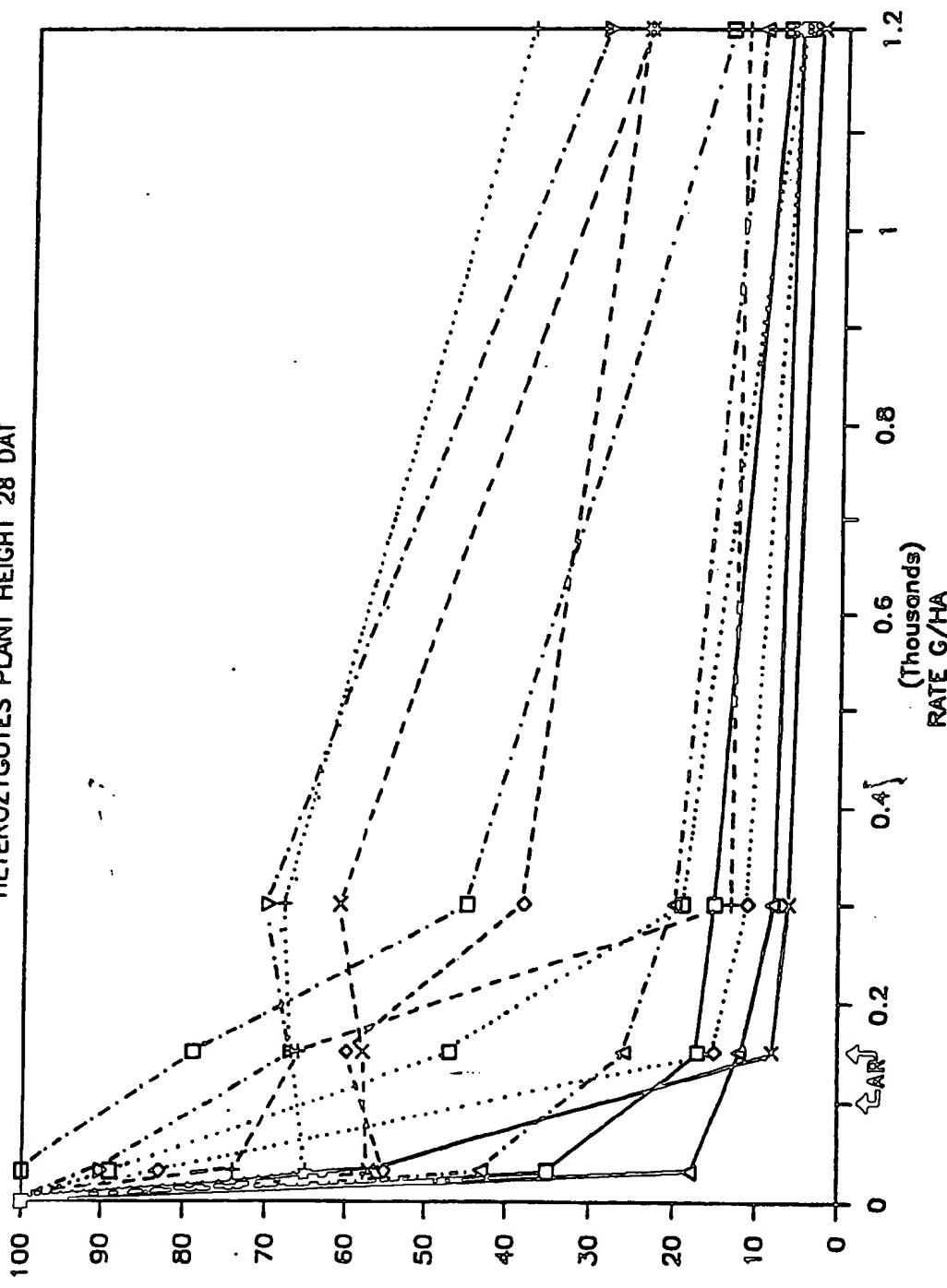
HETEROZYGOTES PLANT HEIGHT 28 DAT



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FIG. 18. SCEPTER DOSE RESPONSE

HETEROZYGOTES PLANT HEIGHT 28 DAT

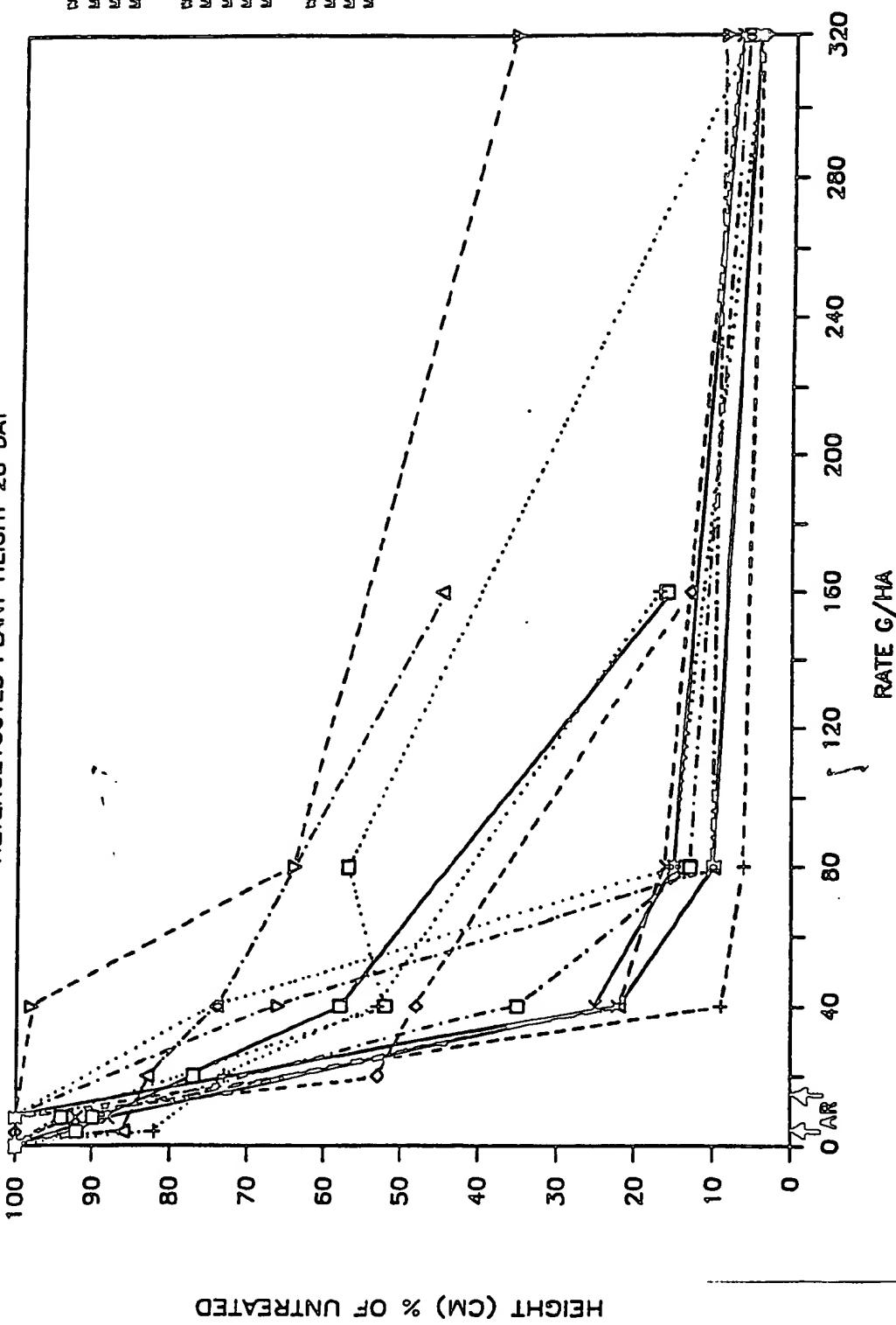


HEIGHT (CM) % OF UNTREATED

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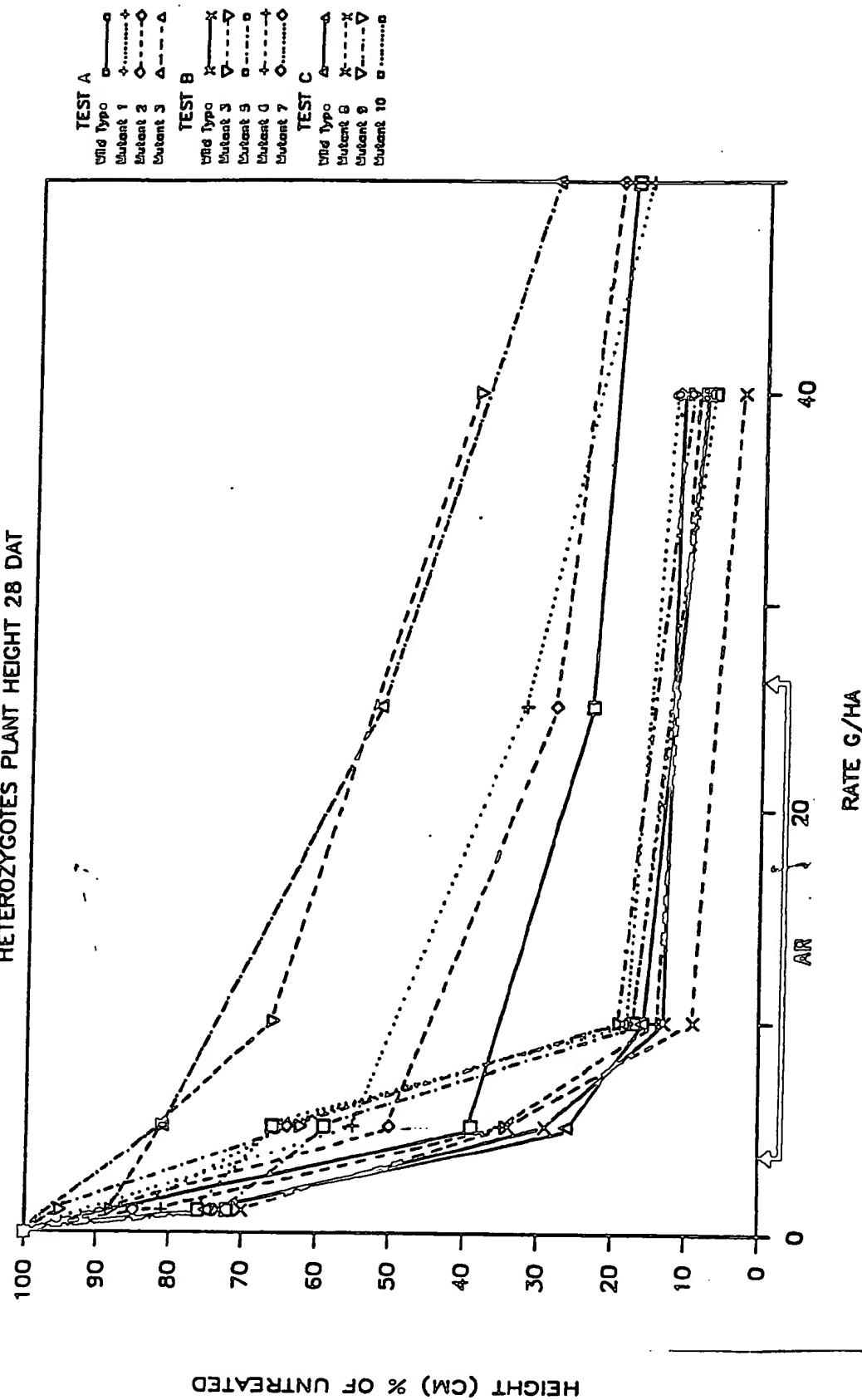
FIG. 19. CLASSIC DOSE RESPONSE

HETEROZYGOTES PLANT HEIGHT 28 DAT

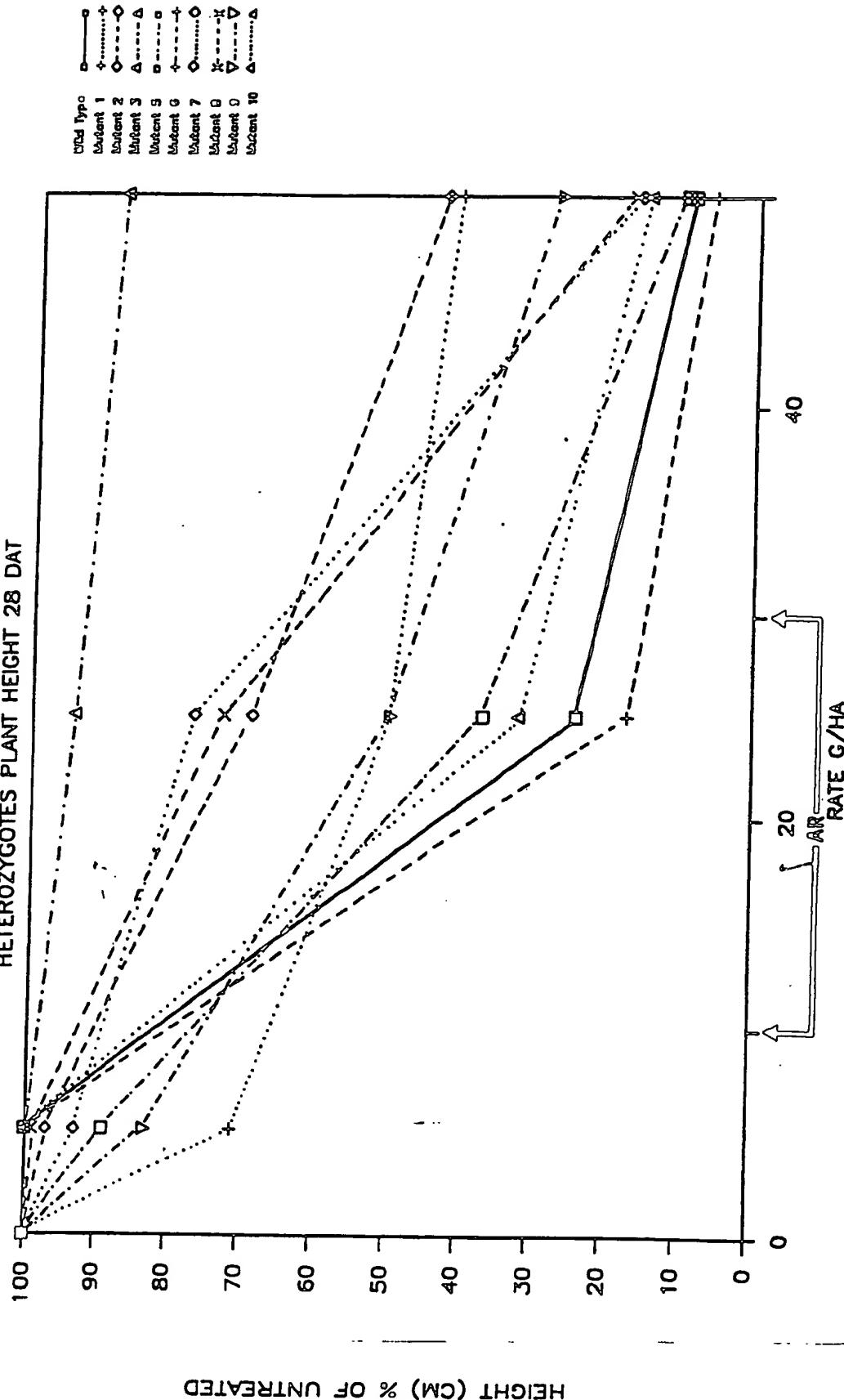


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FIG. 20. GLEAN DOSE RESPONSE
HETEROZYGOTES PLANT HEIGHT 28 DAT

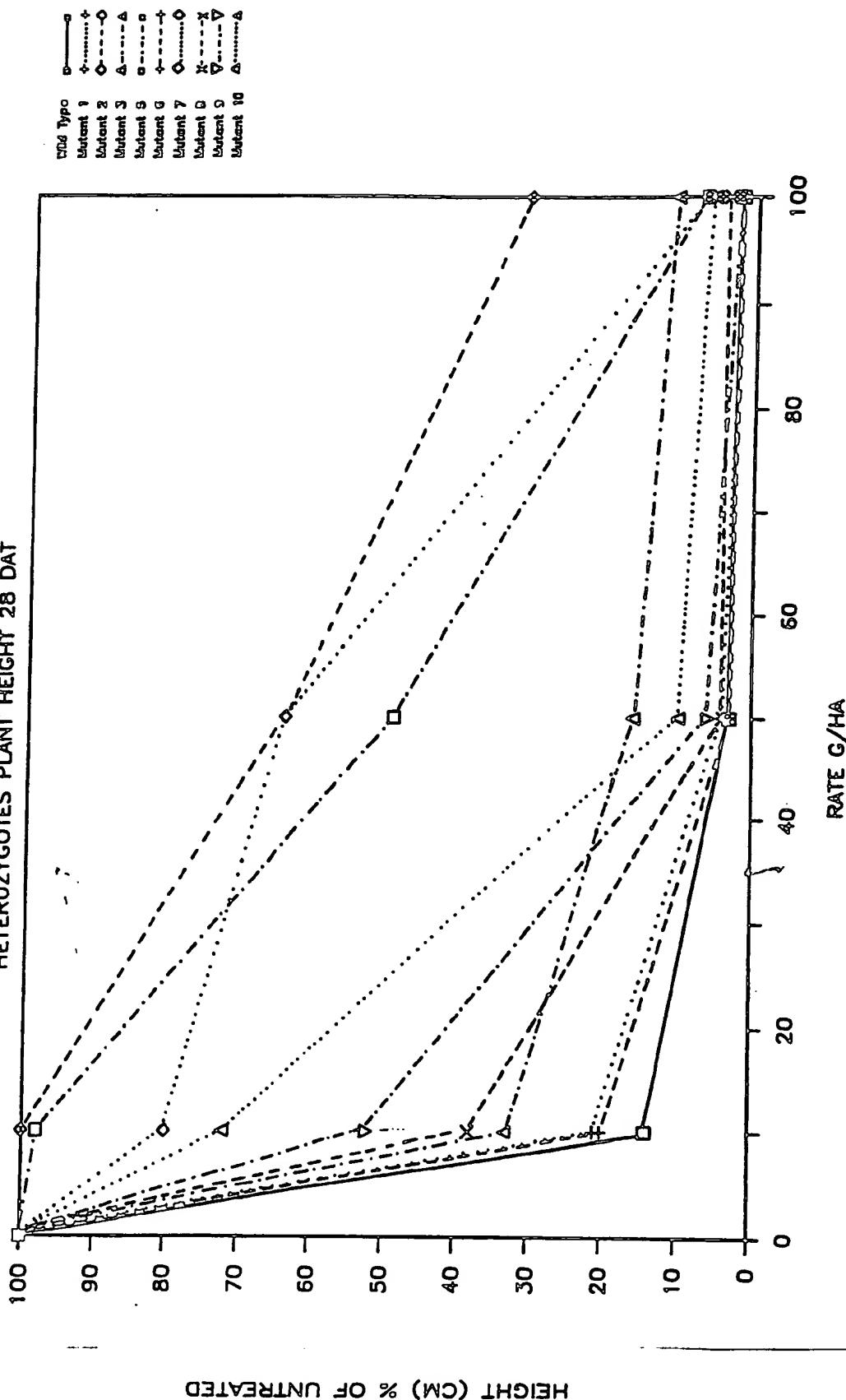


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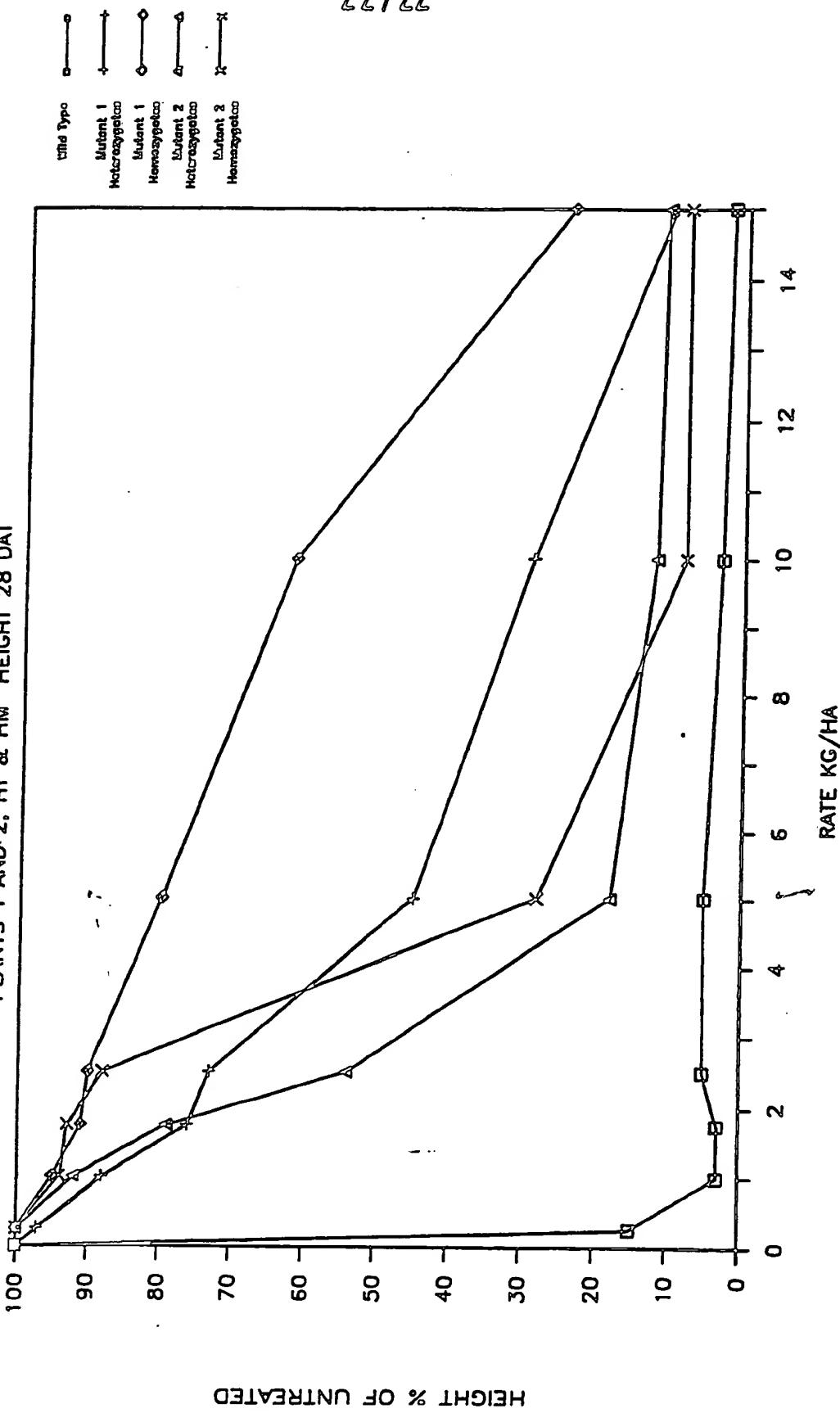
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FIG. 22. PHENOXYPYRIMIDINE DOSE RESPONSE



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FIG. 23. PURSUIT DOSE RESPONSE
PLANTS 1 AND 2. HT & HM HEIGHT 28 DAT



INTERNATIONAL SEARCH REPORT

International Application No. PCT/GB 90/00753

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) ⁸

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC⁵: A 01 H 1/06, C 12 N 15/60

II. FIELDS SEARCHED

Minimum Documentation Searched ⁹

Classification System	Classification Symbols
IPC ⁵	A 01 H, C 12 N

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched ⁸

III. DOCUMENTS CONSIDERED TO BE RELEVANT ¹⁰

Category ¹¹	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
Y	Maydica XXIII, 1978, M.G. Neuffer et al.: "Paraffin oil technique for treating mature corn pollen with chemical mutagens", pages 21-28 see page 21, abstract cited in the application --	1-6
Y	EP, A, 0154204 (MOLECULAR GENETICS, INC.) 11 September 1985 see page 12, lines 15-34; page 13, lines 19-22; page 13, line 31 - page 15, line 30; page 19, lines 7-15, 21-25; page 20, lines 10-21; page 22, line 21 - page 23, line 8; page 49 - page 50, line 4; claims 1, 3-5, 7-9, 24 & US, A, 4761373 (cited in the application)	1-6
X	--	13-14
P, Y	Chemical Abstracts, volume 112, no. 3, 15 January 1990, (Columbus, Ohio, US), ./.	1-6

* Special categories of cited documents: ¹⁰

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"A" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

21st August 1990

Date of Mailing of this International Search Report

25. 04. 90

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

R.J. Eernisse

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

E.B. Swanson et al.: "Microspore mutagenesis and selection: canola plants with field tolerance to the imidazolinones", see page 304, abstract 18959k, & Theor. Appl. Genet. 1989, 78(4), 525-30

V. OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSearchable:

This International Search Report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. Claim numbers 7-12 because they relate to subject matter not required to be searched by this Authority, namely:

See PCT-Rule 39.1(IV): Methods for treatment of the human or animal body by surgery or therapy, as well as diagnostic methods.

2. Claim numbers _____, because they relate to parts of the International application that do not comply with the procedural requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. Claim numbers _____ because they are dependent claims and are not drafted in accordance with the second and third conditions of PCT Rule 8.4(a).

VI. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING:

This International Searching Authority found multiple inventions in this International application as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims of the International application.

2. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims of the International application for which fees were paid, specifically claims:

3. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:

4. As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remarks on Protocol:

- The additional search fees were accompanied by applicant's protocol.
- No protocol accompanied the payment of additional search fees.

ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.

GB 9000753

SA 36844

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 19/09/90. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP-A- 0154204	11-09-85	AU-A-	3950785	12-09-85
		AU-A-	4608989	29-03-90
		JP-A-	60210929	23-10-85
		US-A-	4761373	02-08-88